

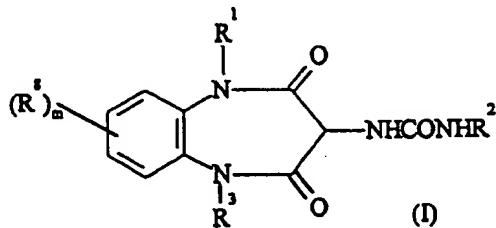
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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : C07D 243/12	A1	(11) International Publication Number: WO 93/14074 (43) International Publication Date: 22 July 1993 (22.07.93)
(21) International Application Number: PCT/EP93/00098		(74) Agents: BREWER, Christopher, Laurence et al.; Glaxo Holdings plc, Glaxo House, Berkeley Avenue, Greenford, Middlesex UB6 0NN (GB).
(22) International Filing Date: 19 January 1993 (19.01.93)		
(30) Priority data: 9201180.8 21 January 1992 (21.01.92) GB		(81) Designated States: AT, AU, BB, BG, BR, CA, CH, CZ, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SK, UA, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, SN, TD, TG).
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(54) Title: 1,5-BENZODIAZEPINE DERIVATIVES AND THEIR USE IN MEDICINE



(57) Abstract

Compounds of general formula (I), wherein R¹ represents a phenyl, C₃-cycloalkyl, C₇₋₁₁ bridged cycloalkyl or C₁₋₆alkyl group which alkyl group may be substituted by a hydroxy, phenyl, C₁₋₆alkoxycarbonyl, C₃-cycloalkyl, or C₇₋₁₁ bridged cycloalkyl group; R² represents a substituted or unsubstituted phenyl group (wherein the substituents may be 1 or 2 of halo, C₁₋₄alkyl, nitro, cyano trifluoromethyl, trifluoromethoxy, C₁₋₄alkylthio or (CH₂)_nR⁴ wherein R⁴ is hydroxy, C₁₋₄alkoxy, CO₂R⁵ or NR⁶R⁷. R³ is phenyl optionally substituted by one or two halogen atoms; R⁵ represents hydrogen or a C₁₋₄alkyl group; R⁶ and R⁷ independently represent hydrogen or a C₁₋₄alkyl group. R⁸ represents hydrogen or a halogen atom; m is 0, 1 or 2; n is 0 or 1; are antagonists of gastrin and CCK-B receptors.

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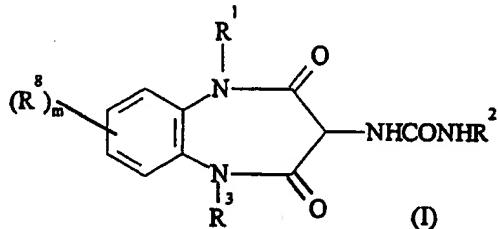
1,5 - BENZODIAZEPINE DERIVATIVES AND THEIR USE IN MEDICINE

5 This invention relates to novel 1,5-benzodiazepine derivatives, to processes for their preparation, to pharmaceutical compositions containing them and to their use in medicine.

In particular the invention relates to compounds which are potent and specific antagonists of gastrin and/or cholecystokinin (CCK).

Thus, the invention provides compounds of general formula (I)

10



15

wherein

R¹ represents a phenyl, C₃-cycloalkyl, C_{7,11} bridgedcycloalkyl or C_{1,6}alkyl group which alkyl group may be substituted by a hydroxy, phenyl, C_{1,6}alkoxycarbonyl, C₃-cycloalkyl, or C_{7,11} bridgedcycloalkyl group;

20 R² represents a substituted or unsubstituted phenyl group (wherein the substitutents may be 1 or 2 of halo, C_{1,4}alkyl, nitro, cyano, trifluoromethyl, trifluoromethoxy, C_{1,4}alkylthio or (CH₂)_nR⁴ wherein R⁴ is hydroxy, C_{1,4}alkoxy, CO₂R⁵ or NR⁶R⁷.

R³ is phenyl optionally substituted by one or two halogen atoms;

R⁵ represents hydrogen or a C_{1,4}alkyl group;

R⁶ and R⁷ independently represent hydrogen or a C_{1,4}alkyl group.

25 R⁸ represents hydrogen or a halogen atom; m is zero, 1 or 2;

n is zero or 1; and pharmaceutically acceptable salts and solvates thereof.

It will be appreciated that compounds of formula (I) possess at least one asymmetric carbon atom (namely the carbon atom occupying the 3-position of the diazepine ring) and the compounds of the invention thus include all stereoisomers and mixtures thereof including the racemates.

30

In the compounds of formula (I) 'alkyl' when used as a substituent or part of a substituent group means that the group may be straight or branched. Thus, C₁₋₆ alkyl includes methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl or tert-butyl, n-pentyl, isopentyl neopentyl, n-hexyl, iso hexyl, 1,3-dimethylbutyl, 3,3-dimethylbutyl,
5 2,3-dimethylbutyl.

For the group R¹ the term C₃₋₇cycloalkyl as a group or part of a group refers to a monocyclic alkyl group such as cyclopropyl, cylobutyl, cyclopentyl, cyclohexyl or cycloheptyl. The term C₇₋₁₁ bridged cycloalkyl refers to groups such adamantly, norbornanyl or norbornenyl.

10 For the groups R⁵ R⁶ and R⁷ the term C₁₋₄alkyl includes 3-4-cycloalkyl (e.g. cyclopropyl or cyclobutyl) as well as straight or branched chain alkyl groups as defined above.

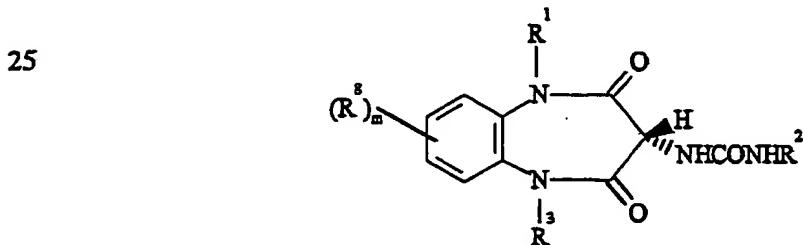
Halogen in the definition of compounds of formula (I) may represent a fluoro, chloro, bromo or iodo substituent.

15 When R² is a phenyl group substituted by a single substituent this may be in the ortho, para or more preferably in the meta position.

When R⁸ is halogen this is preferably chlorine or fluorine.

When m is 1 or 2 the halogen atom(s) e.g. chlorine or fluorine are preferably in the 7 and/or 8 positions.

20 The compounds of formula (I) posses at least one asymmetric carbon atom (namely the carbon atom occupying the 3-position of the diazeine ring) and particularly preferred compounds of the invention or those having the relative stereochemistry shown in formula (1a)



wherein the solid wedge bond indicates the group is above the plane of the diazepine ring and the broken bond indicates the group is below the plane of the diazepine ring.

When R¹ represents an alkyl group substituted by a hydroxyl group this is preferably a C₃₋₆alkyl group substituted by hydroxy. Examples of such groups include

- 5 2-hydroxypropyl, 2-hydroxy-3-methylbutyl and 2-hydroxy-3,3-dimethylbutyl of which 2-hydroxy-3-methylbutyl, and 2-hydroxy-3,3-dimethylbutyl are particularly preferred.

When R¹ represent an alkyl group substituted by a C₃₋₇cycloalkyl group this is preferably a C₂₋₃alkyl group such as ethyl or 1-methylethyl, substituted by a C₃₋₇cycloalkyl group such as cyclopentyl.

- 10 When R¹ is a bridged C₇₋₁₁cycloalkyl group this may be for example an adamantyl group such as 1-adamantyl or 2-adamantyl group or a 2-norbornanyl group.

- When R¹ is an alkyl group substituted by a bridged C₇₋₁₁cycloalkyl group this is preferably an ethyl group or more especially a methyl group substituted by a bridged C₇₋₁₁cycloalkyl group. Examples of suitable brigid cycloalkyl groups include adamantyl such as 1-adamantyl or 2-adamantyl, 2-norbornanyl or 5-norbornenyl. Most preferably R¹ represents 1-adamantylmethyl.

When R¹ is alkyl substituted by phenyl this may be for example benzyl or phenethyl.

When R¹ is alkyl substituted by alkoxy carbonyl this is preferably methyl substituted by alkoxy carbonyl such methoxycarbonyl or as t-butoxycarbonyl.

- 20 A preferred class of compounds of formula (I) is that in which R¹ represents a phenyl, adamantyl, norbornanyl, phenethyl, C₄₋₆alkyl e.g. n- butyl, 3-methyl butyl, 3,3-dimethyl butyl, 1,3-dimethylbutyl, 2,3- dimethylbutyl, C₃₋₆ hydroxy alkyl e.g. 2-hydroxypropyl, 2-hydroxy-3- methylbutyl, 2-hydroxy-3,3-dimethylbutyl, C₁₋₂alkyl substituted by a bridged C₇₋₁₀cycloalkyl group e.g. 2-norbornylmethyl, 5-norbornenylmethyl, 2-adamantylmethyl, 2-adamantylethyl, 2-(1-adamantyl)ethyl, 1-adamantylmethyl, alkoxy carbonylalkyl, e.g. methoxycarbonylmethyl or t-butoxycarbonylmethyl, or 2-cyclopentylethyl.

A particularly preferred class of compounds of formula (I) is that in which R¹ is 3-methylbutyl, 3,3-dimethylbutyl, 2-hydroxy-3-methylbutyl, 2-hydroxy-3,3-dimethylbutyl, 2-cylopentylethyl, 5-norbornenylmethyl or 1-adamantylmethyl.

A further preferred class of compounds of formula (I) is that in which R² represents phenyl optionally substituted by bromine, chlorine, fluorine, methyl, methoxy, methylthio, trifluoromethoxy, cyano, dimethylamino or (CH₂)_nCO₂R⁵ wherein R⁵ is hydrogen or ethyl. Most preferably R² represents phenyl optionally substituted by methoxy, dimethylamino, cyano, methylthio, CO₂H or CO₂C₂H₅.

5 A further preferred class of compounds of formula (I) is that in which R³ represents phenyl or phenyl mono- or di-substituted by fluorine, preferably in the ortho and/or para position(s). Preferably R³ represents unsubstituted phenyl or orthofluorophenyl.

10 A preferred group of compounds of formula (I) those wherein R¹ represents C₄₋₆alkyl such as 3-methylbutyl, 3,3-dimethylbutyl, 2-hydroxy-3-methylbutyl, 2-hydroxy-3,3-dimethylbutyl 2-cyclopentylethyl, 5-norbornenylmethyl or 1 adamantlylmethyl; R² represents phenyl or phenyl substituted by methoxy, cyano, nitro, carboxyl, ethoxycarbonyl, methylthio, or dimethylamino and preferably the substituent is in the meta 1-position;

15 R³ represents phenyl or ortho fluorophenyl; R⁸ represents hydrogen, chlorine or fluorine; and enantiomers and salts thereof.

20 A particularly preferred group of compounds of formula (I) are those wherein R¹ is 3-methylbutyl; R² is phenyl optionally substituted in the meta position by methylthio or dimethylamino group; R³ is phenyl or ortho fluorophenyl; R₈ is hydrogen or chlorine or fluorine and m is zero, 1 or 2.

25 A further particularly preferred group of compounds of formula (I) are those wherein R¹ represents 1-adamantlylmethyl R² is phenyl optionally substituted in the meta position by a methyl, methoxy, methylthio, nitro, dimethylamino, ethoxycarbonyl or carboxyl group; R³ is phenyl and R⁸ is hydrogen. Within this group especially preferred compounds are those wherein R² is phenyl optionally substituted by dimethylamino, ethoxycarbonyl or carboxyl group

Preferred compounds according to the invention include:

N-phenyl-N-[2,3,4,5-tetrahydro-2,4-dioxo-1-(3-methylbutyl)-5-phenyl-1H-1,5-benzodiazepin-3-yl]urea;

- N-[1-(3,3-Dimethyl-2-hydroxybut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-
3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea
N-phenyl-N'-[2,3,4,5-tetrahydro-2,4-dioxo-1-(3,3-dimethylbutyl)-5-phenyl-1H-1,5-
benzodiazepin-3-yl]urea;
- 5 N-phenyl-N'-[2,3,4,5-tetrahydro-2,4-dioxo-1-(1-adamantylmethyl)-5-
phenyl-1H-1,5-benzodiazepin-3-yl]urea;
N-[2,4-Dioxo-1-(2-hydroxy-3-methylbutyl)-5-phenyl-2,3,4,5-
tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea
N-(3-dimethylaminophenyl)-N'-[2,3,4,5,-tetrahydro-2,4-dioxo-1-(3-
10 methylbutyl)-5-(2-fluorophenyl)-1H-1,5-benzodiazepin-3-yl]urea;
N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-
benzodiazepin-3- yl]-N'-(3-ethoxycarbonylphenyl)urea
N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-
benzodiazepin-3- yl]-N'-(3-(N,N-dimethylamino)phenyl)urea
- 15 N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-
benzodiazepin-3- yl]-N'-(3-carboxyphenyl)urea
N-[1-(Adamantane-1-methyl)-2,4-dioxo-7-fluoro-5(4-fluorophenyl)-2,3,4,5-tetrahydro-
1H-1,5benzodiazepin-3-yl]-N'(3- dimethylamino)phenylurea
and (+) enantiomers and salts thereof.
- 20 Particularly preferred compounds according to the invention include:
N-phenyl-N'-[2,3,4,5-tetrahydro-2,4-dioxo-1-(1-adamantylmethyl)-5-
phenyl-1H-1,5-benzodiazepin-3-yl]urea;
N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-
benzodiazepin- 3-yl]-N'-(3-carboxyphenyl)urea
25 N-phenyl-N'-[2,3,4,5-tetrahydro-2,4-dioxo-1-(3-methylbutyl)-5-
phenyl-1H-1,5-benzodiazepin-3-yl]urea;
N-(3-dimethylaminophenyl)-N'-[2,3,4,5,-tetrahydro-2,4-dioxo-1-(3-
methylbutyl)-5-(2-fluorophenyl)-1H-1,5-benzodiazepin-3-yl]urea;
and the (+) enantiomers thereof and salts thereof.

The pharmaceutically acceptable salts of the compounds of formula (I) include conventional salts formed for example from pharmaceutically acceptable inorganic or organic acids as well as quaternary ammonium acid addition salts. Examples of suitable salts include hydrochloric, hydrobromic, sulphuric, phosphoric, nitric, perchloric, fumaric, 5 acetic, propionic, succinic, glycolic, formic, lactic, maleic, tartaric, citric, pamoic, malonic, hydroxymaleic, phenylacetic, glutamic, benzoic, salicylic, fumaric, toluenesulphonic, methanesulphonic, naphthalene-2-sulphonic, benzenesulphonic and the like. Other acids such as oxalic, while not in themselves pharmaceutically acceptable, may be useful in the preparation of salts useful as intermediates in obtaining the compounds of 10 the invention and their pharmaceutically acceptable salts.

The compounds of formula (I) in which R⁵ represents hydrogen may form pharmaceutically acceptable salts with suitable cations. Suitable pharmaceutically acceptable cations include alkali metal (e.g. sodium or potassium) and alkaline earth metal (e.g calcium or magnesium) cations.

15 References hereinafter to a compound according to the invention includes both compounds of formula (I) and their pharmaceutically acceptable salts and solvates.

The compounds of the invention are potent and specific antagonists of gastrin and/or CCK. The compounds of the invention have been shown to be antagonists of CCK, particularly at CCK-B receptors as demonstrated for example by the compound's ability 20 to inhibit the contractile actions of CCK-4 in the presence of a CCK-A antagonist, in the guinea-pig isolated ileum longitudinal muscle- myenteric plexus.

The compounds of the invention have also been shown to be antagonists of gastrin as demonstrated by their ability to inhibit pentagastrin-stimulated acid secretion from rat isolated gastric mucosa using the procedure described by J.I. Reeves and R. Stables in Br. 25 J. Pharmac., 1985, 86, p.677-684.

Compounds of the invention have also been found to have a significantly weaker activity at CCK-A receptors compared with their activity at gastrin and/or CCK-B receptors, as demonstrated by their ability to inhibit the contractile activity of CCK-8 in guinea-pig siolated ileum longitudinal muscle-myenteric plexus.

The preparation and use of guinea-pig isolated ileum longitudinal muscle-myenteric plexus has been described by K-H Buchheit et al in Nauyn-Schmeideberg's Arch. Pharmacol, (1985), 329, p36-41 and by V.L. Lucaites et al (1991) in J. Pharmacol. Exp. Ther., 256, 695-703.

5 The greater affinity of the compounds of the invention for the CCK-B receptor over the CCK-A receptor has also been established using the CCK receptor binding assays described by G Dal Fornos et al., J. Pharmacol. Exp & Ther. 261, 1056-1063, 1992.

10 The compounds of the invention are therefore useful for the treatment and/or prevention of disorders in mammals, especially humans, where modification of the effects of gastrin or CCK is of therapeutic benefit. Thus the compounds of the invention are useful for the treatment of central nervous system disorders where CCK and/or gastrin are involved. For example anxiety disorders (including panic disorder, agoraphobia, social phobia, simple phobia, obsessive compulsive disorders, post traumatic stress disorder, and general anxiety disorder), tardive dyskinesia, depression, Parkinson's disease or psychosis.

15 The compounds of the invention are also useful for the treatment of gastrointestinal disorders especially those where there is an advantage in lowering gastric acidity. Such disorders include peptic ulceration, reflux oesophagitis and Zollinger Ellison syndrome. They may also be useful for the treatment of gastrointestinal disorders such as irritable bowel syndrome, excess pancreatic secretion, acute pancreatitis, motility disorders, antral

20 G cell hyperplasia, fundic mucosal hyperplasia or gastrointestinal neoplasms. They may also be useful for the treatment of dependency on drugs or substances of abuse and withdrawal, Gilles de la Tourette syndrome, or dysfunction of appetite regulatory systems; as well as the treatment of certain tumours of the lower oesophagus, stomach, intestines and colon. Compounds of the invention are also useful for directly inducing

25 analgesia, or enhancing opiate or non-opiate mediated analgesia, as well as anaesthesia or loss of the sensation of pain.

Compounds of the invention have also been found to exhibit anxiolytic activity in conventional pharmacological tests. For example in mice in the black-white box test and in the rat social interaction model.

The invention therefore provides a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof for use in therapy, in particular in human medicine.

According to another aspect the invention provides the use of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof for the manufacture of 5 a medicament for the treatment of conditions where modification of the effects of gastrin and/or CCK is of therapeutic benefit.

According to a further aspect of the invention we provide a method for the treatment of a mammal, including man, in particular in the treatment of conditions where modification of the effects of gastrin and/or CCK is of therapeutic benefit which method 10 comprises administering an effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof to the patient.

It will be appreciated by those skilled in the art that reference herein to treatment extends to prophylaxis as well as the treatment of established diseases or symptoms.

It will further be appreciated that the amount of a compound of the invention 15 required for use in treatment will vary with the nature of the condition being treated and the age and the condition of the patient and will be ultimately at the discretion of the attendant physician or veterinarian. In general however doses employed for adult human treatment will typically be in the range of 0.01-2000mg per day e.g 0.01-500mg per day.

The desired dose may conveniently be presented in a single dose or as divided doses 20 administered at appropriate intervals, for example as two, three, four or more sub-doses per day.

Because the compounds of the invention antagonise the function of CCK in animals, they may also be used as feed additives to increase the food intake in animals in daily dosages of around 1mg/kg to 10mg/kg.

25 While it is possible that, for use in therapy, a compound of the invention may be administered as the raw chemical it is preferable to present the active ingredient as a pharmaceutical formulation.

The invention thus further provides a pharmaceutical formulation comprising a compound of formula (I) or a pharmaceutically acceptable salt thereof together with one 30 or more pharmaceutically acceptable carriers therefor and, optionally, other therapeutic

and/or prophylactic ingredients. The carrier(s) must be 'acceptable' in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof.

5 The compositions of the invention include those in a form especially formulated for oral, buccal, parenteral, implant, or rectal administration. Oral administration is preferred.

Tablets and capsules for oral administration may contain conventional excipients such as binding agents, for example, syrup, accacia, gelatin, sorbitol, tragacanth, hydroxypropyl cellulose, mucilage of starch or polyvinylpyrrolidone; fillers, for example, lactose, sugar, microcrystalline cellulose, maize-starch, calcium phosphate or sorbitol; 10 lubricants, for example, hydrogenated vegetable oils, magnesium stearate, stearic acid, talc, polyethylene glycol or silica; disintegrants, for example, potato starch or sodium starch glycollate, or wetting agents such as sodium lauryl sulphate. The tablets may be coated according to methods well known in the art. Oral liquid preparations may be in the form of, for example, aqueous or oily suspensions, solutions emulsions, syrups or 15 elixirs, or may be presented as a dry product for constitution with water or other suitable vehicle before use. Such liquid preparations may contain conventional additives such as suspending agents, for example, sorbitol syrup, methyl cellulose, glucose/sugar syrup, gelatin, hydroxyethylcellulose, carboxymethyl cellulose, aluminium stearate gel or hydrogenated edible fats; emulsifying agents, for example, lecithin, sorbitan mono-oleate 20 or acacia; non-aqueous vehicles (which may include edible oils), for example, almond oil, fractionated coconut oil, oily esters, propylene glycol or ethyl alcohol; and preservatives, for example, methyl or propyl p-hydroxybenzoates or sorbic acid. The compositions may also be formulated as suppositories, e.g. containing conventional suppository bases such as cocoa butter or other glycerides.

25 For buccal administration the composition may take the form of tablets or lozenges formulated in conventional manner.

The composition according to the invention may be formulated for parenteral administration by injection or continuous infusion. Formulations for injection may be 30 presented in unit dose form in prefilled syringes, vials and ampoules, or in multi-dose containers with an added preservative. The compositions may take such forms as

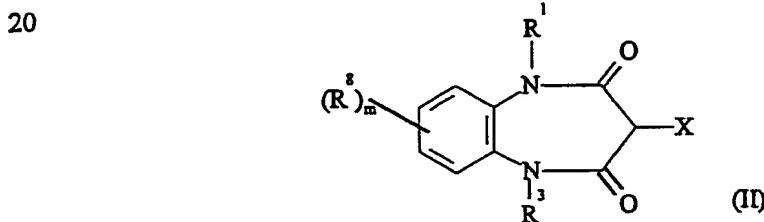
suspensions, solutions, or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilising and/or dispersing agents. Alternatively the active ingredient may be in powder form which may be obtained by freeze drying for constitution with a suitable vehicle, e.g. sterile, pyrogen-free water, before use.

5 The composition according to the invention may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection. Thus for example, the compounds of the invention may be formulated with suitable polymeric or hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

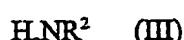
10 The compositions according to the invention may contain between 0.1 - 99% of the active ingredient, conveniently from 30-95% for tablets and capsules and 3-50% for liquid preparations.

15 Compounds of general formula (I) and salts thereof may be prepared by the general methods outlined hereinafter. In the following description, the groups R¹-R⁸ are as defined for the compounds of formula (I) unless otherwise stated.

20 According to a first general process (A) compounds of formula (I) may be prepared by reacting a compound of formula (II) in which X represents the group -N=C=O, or NHCOR⁹ wherein R⁹ is an optionally substituted phenoxy group or a 1-imidazole group.



25 with an amine of formula (III)

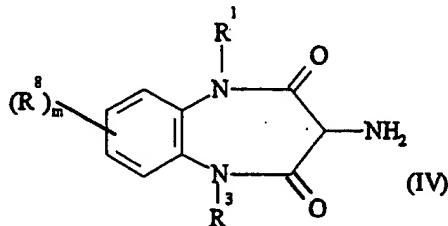


30 optionally in the presence of a base such as a tertiary amine (e.g. triethylamine). The reaction conveniently takes place in a suitable solvent such as a halogenated hydrocarbon

(e.g. dichloromethane) or an ether (e.g. tetrahydrofuran) or an amide e.g. N,N-dimethylformamide optionally at a temperature ranging from room temperature to the reflux temperature of the solvent.

In a particular aspect of the process (A) when X is the group NHCOR^9 and R^9 is a 5 1-imidazole group, the imidazolide (II) may be formed *in situ* in which case the amine of formula (III) will be mixed with a compound of formula (IV)

10



in the presence of carbonyldiimidazole under the aforementioned conditions.

For process A when X is the group NHCOR^9 and R^9 is optionally substituted phenoxy group the reaction with the primary amine (III) is preferably carried out in the 15 presence of a base such as a tertiary amine e.g. triethylamine.

For process A when X is the isocyanate group -N=C=O the reaction with the primary amine (III) is preferably carried out in an aprotic solvent such as a halohydrocarbon e.g. methylene chloride. Conveniently the isocyanate is generated *in situ* prior to the addition of the primary amine (III).

20 The compounds of formula (II) wherein R^9 is an optionally substituted phenoxy group may be prepared from the primary amine (IV) by reaction with the corresponding optionally substituted phenyl chloroformate in the presence of a base such as pyridine. The reaction may be carried out in a solvent such as a halohydrocarbon e.g. dichloromethane and at a temperature from 0-50°.

25 Compounds of formula (II) wherein R^9 is a 1-imidazole group may be prepared by reacting a compound of formula (IV) with carbonyldiimidazole in the presence of a suitable solvent such as a halogenated hydrocarbon (e.g. dichloromethane) or an ether (e.g. tetrahydrofuran) at a temperature ranging from 0° to 80° (conveniently at room temperature).

Compounds of formula (II) wherein X is the isocyanate grouping -N=C=O may be prepared from the primary amine (IV) by reaction with phosgene (COCl₂) in a suitable solvent such as methylene chloride.

According to a further general process (B) compounds of formula (I) may be
5 prepared by reacting a compound of formula (IV)
with an isocyanate of formula (V)



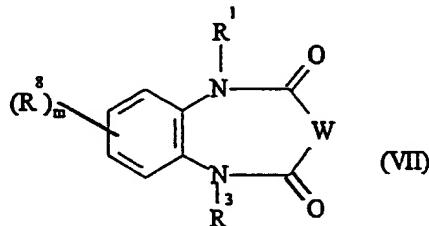
10 or a carbamoyl chloride of formula (VI)



The reaction conveniently takes place in the presence of a suitable solvent such as a
15 halohydrocarbon (e.g. dichloromethane), an ether (e.g tetrahydrofuran) or a nitrile (e.g. acetonitrile) or a mixture thereof at a temperature in the range of 0°C to 80°C.

Compounds of formula (IV) may be prepared by reduction of compounds of formula
(VII)

20



wherein W is CH-N₃ or C=N-NHPh.

25

Compounds of formula (VII) wherein W is CH-N₃ may be reduced to a compound of formula (IV) by hydrogenation in the presence of a suitable catalyst such as palladium, on a support such as carbon or calcium carbonate, or platinum (IV) oxide. The reaction conveniently takes place in the presence of a solvent such as an alkanol (e.g. ethanol) an ester (e.g. ethyl acetate) or acetic acid.

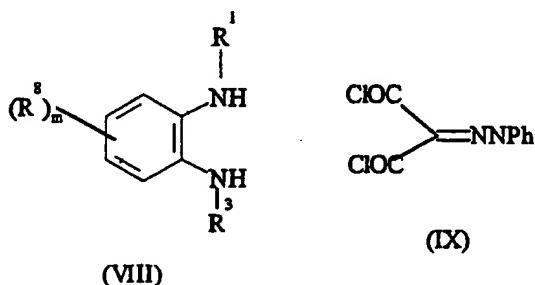
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Compounds of formula (VII) wherein W is C=N-NHPh may be reduced to a compound of formula (IV) by reaction with zinc and acetic acid. This reaction may be carried out a temperature with the range 0-50°.

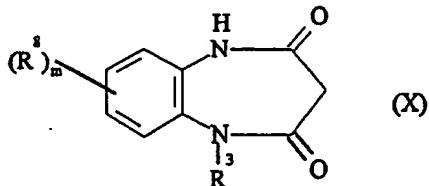
Compounds of formula (VII) wherein W is CHN, may be prepared from a compound of formula (VII) wherein W is CH₂ by treatment with a strong base such as sodium hydride or potassium tert-butoxide followed by tri-isopropyl benzenesulphonyl azide. The reaction conveniently takes place in a solvent such as an ether (e.g. tetrahydrofuran) at a temperature in the range of -78° to 20°.

Compounds of formula (VII) in which W is C=NNHPh may be prepared by reaction of the ortho-phenylenediamine (VIII) with the diacid chloride (IX), in a suitable solvent such as an ether e.g. tetrahydorfuran

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with a compound R¹Y where Y is halogen (e.g.a chlorine or bromine atom) or a mesylate group under strongly basic conditions. Thus the reaction may conveniently be carried out by pretreating the compound of formula (X) with a strong base such as sodium hydride in a suitable aprotic solvent such as an amide (e.g. N,N- dimethylformamide) at a temperature ranging from 0° to reflux.

In the above described reaction scheme when the group R¹ contains an hydroxyl group then this may be present in a protected form e.g. as an ether such as an arylmethyl ether e.g. a benzyl ether.

Compounds of formula (VIII) are either known compounds or may be prepared by 5 analogous methods. Thus for example a compound of formula (VIII) may be prepared by alkylation of the amine (XI).



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Thus the amine (XI) may be reacted with the compound R¹Y, in which Y is chlorine or bromine, optionally in the presence of sodium iodide in a solvent such as N,N-dimethylformamide.

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Compounds of formula (VIII) wherein R¹ represents the group -CH₂-CH(OH)R¹, where R¹ is a C₁₋₄alkyl group may be prepared by reaction of compound (XI) with the epoxide (XII) in a solvent such as an alkanol e.g. ethanol and in the presence of an acid catalyst such as p-toluene sulphonic acid.

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Compounds of formula (VIII) where in R¹ is an optionally substituted alkyl group. May also be prepared from compound (XI) by reaction with a suitable aldehyde or ketone with concomitant or subsequent reduction of the reaction product. Thus for example a compound formula (VIII) wherein R¹ is 1,3-dimethylbutyl may be prepared from compound (XII) by reaction with methylisobutyl ketone followed by reaction with sodium borohydride.

In general, the compounds of formula (III), V and (VI) are either known compounds or may be prepared according to methods used for the preparation of known compounds,

According to a further process (C) a compound of formula (I) may be converted into another compound of formula (I) using conventional techniques.

Thus compounds of formula (I) wherein R² is a phenyl group substituted by a carboxyl group may be prepared by hydrolysis of the corresponding compound of formula (I) wherein R² is a phenyl group substituted by an alkoxy carbonyl group.

In the processes described above the group R¹ and R² in the intermediates II, III, V and VI may be a group as defined in formula (I) or a group convertible thereto.

The foregoing series of reactions involve a number of alternative pathways which may start with the 1,5-benzodiazepine of formula (X) as defined above. Thus according to a further general process (D) a compound of formula (I) may be prepared by reacting a compound of formula (X) in one or more stages with reagents serving to introduce the groups R¹ and NHCONHR².

Compounds of formula (I) contain at least one asymmetric carbon atom, namely the carbon atom of the diazepine ring to which the substituted urea grouping is attached. Specific enantiomers of the compounds of formula (I) may be obtained by resolution of the racemic compound using conventional procedures such as chiral HPLC. Alternatively the required enantiomer may be prepared by the corresponding enantiomeric amine of formula (IV) using any of the processes described above for preparing compounds of formula (I) from the amine (IV). The enantiomers of the amine (IV) may be prepared from the racemic amine (IV) using conventional procedures such as salt formation with a suitably optically active acid such as R- camphorsulphonic acid.

The following examples, which are non-limiting, illustrate the invention.

In the Preparations and Examples, unless otherwise stated: Melting points (m.p.) were determined on a Buchi m.p. apparatus and are uncorrected. All temperatures refer to 0C. Infrared spectra were measured in chloroform-d₁ solutions on a FT-IR instrument. Proton Magnetic Resonance (1H-NMR) spectra were recorded at 300MHz as solutions in chloroform-d₁. Chemical shifts are reported in ppm downfield (δ) from Me4Si as an internal standard, and are assigned as singlets (s), doublets (d), doublet of doublets (dd) or multiplets (m). Column chromatography was carried out over silica gel (Merck AG Darmstadt, Germany). Solutions were dried over anhydrous sodium sulphate. "Petrol" refers to petroleum ether, b.p. 40-60°C. Dichloromethane was redistilled over calcium hydride; tetrahydrofuran was redistilled over sodium; ethyl ether was redistilled over

sodium and ethyl acetate was dried over activated molecular sieves. The following abbreviations are used in the text. EA = ethyl acetate, CH = cyclohexane, P = petroleum ether 40-60°C, THF = tetrahydrofuran, DCM = dichloromethane, EE = ethyl ether, DMF = N,N-dimethylformamide. Tlc refers to thin layer chromatography on silica plates. All 5 the compounds are intended as racemic mixtures unless otherwise indicated.

Intermediate 1

2-Fluoro-2'-(3-methylbut-1-yl)amino-diphenylamine

10 1-Bromo-3-methylbutane (4.33ml) was added to a solution of the 2-amino-2'-fluorodiphenylamino (7.0g) and sodium iodide (5.24g) in dimethylformamide (250ml) under a nitrogen atmosphere. The solution was stirred at 120° for 8h, then cooled to room temperature, diluted with water (300ml) and extracted with diethyl ether (2x250ml). The combined organic extracts were washed with brine (300ml), dried and 15 concentrated in vacuo to an oil, which was purified by flash chromatography (eluting with CH-EA 95:5) to give the title compound as a yellow oil (6.3g). T.l.c. CH-EA (9:1) Rf 0.75.

Intermediate 2

20 2,4-Dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)-3-phenylhydrazone-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 1 (6.3g) and the 2-phenylhydrazonomalonyl dichloride (6.8g) were each taken up in THF (150ml) and dropped in a flask containing THF (200ml) maintained at -5° under a nitrogen atmosphere. After complete addition the solution was allowed to 25 warm to room temperature and then heated to 50° for 2h. The solution was concentrated in vacuo to an oil, which was purified by flash chromatography (eluting with CH-EA 8:2) to give the title compound as a yellow solid (5.8g). M.p. 104-105° T.l.c. CH-EA (7:3), Rf 0.59.

30 Intermediate 3

3-Amino-2,4-dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 2 (5.8g) in glacial acetic acid (50ml) was added dropwise, to a suspension of zinc dust (6.37g) in glacial acetic acid (20ml) cooled to 0°. The mixture was stirred at 23° for 3h, then diluted with water (200ml) and decanted from zinc. Solid sodium carbonate was added until pH=9 and the mixture extracted with ethyl acetate (2x300ml). The combined organic extracts were washed with brine (300ml), dried and concentrated in vacuo to an oil which was purified by flash chromatography (eluting in gradient from CH-EA 2:1 to EA) to give the title compound as a white foam (2.8g). M.p.

10 125-6°. T.l.c. DCM-methanol (30:1), Rf 0.38.

Intermediate 4

2-(3,3-Dimethylbut-1-yl)amino-2'-fluoro-diphenylamine

Sodium borohydride (22.7g) was added portionwise to a mixture of the 15 2-amino-2'-fluorodiphenylamine (8.0g), sodium acetate trihydrate (16.33g) and 3,3-dimethylbutyraldehyde (5ml) in acetic acid (12.8ml), water (50ml) and ethanol (40ml) cooled to 0°. The solution was stirred at 23° for 30 min., then diluted with ethyl acetate (300ml). The organic layer was washed with a 10% solution of sodium hydroxide (3x200ml) and brine (200ml), dried and concentrated in vacuo to an oil, which was 20 purified by flash chromatography (eluting with CH-EA 9:1) to give the title compound as a yellow oil (7.44g). T.l.c. CH-EA (9:1), Rf 0.85.

Intermediate 5

1-(3,3-Dimethylbut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-3-phenylhydrazone-

25 2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 4 (7.73g) and the 2-phenylhydrazone- malonyldichloride (7.97g) were each taken up in THF (100ml) and dropped in a flask containing THF (300ml) maintained at -5° under a nitrogen atmosphere. After complete addition the solution was allowed to warm to room temperature and then heated to 50° for 3h. The solution was concentrated in vacuo to an oil, which was purified by flash chromatography (eluting with CH-EA 8:2)

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to give the title compound as yellow solid (10.8g). M.p.112-114°. T.l.c. CH-EA (8:2),

Rf 0.40..

Intermediate 6

5 3-Amino-1-(3,3-dimethylbut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 5 (10.1g) in glacial acetic acid (80ml) was added, dropwise, to a suspension of zinc dust (10.8g) in glacial acetic acid (20ml) cooled to 0°. The mixture was stirred at 23° for 2h, then diluted with water (200ml) and decanted from zinc.

10 Solid sodium carbonate was added until pH=9 and the solution then extracted with ethyl acetate (3x250ml). The combined organic extracts were washed with brine (400ml), dried and concentrated in vacuo to an oil, which was purified by flash chromatography (eluting in gradient from CH-EA 2:1 to ethyl acetate) to give the title compound as a white foam (5.4g). M.p. 98-100°. T.l.c. DCM-methanol (20:0.5), Rf 0.3.

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Intermediate 7

2,4-Dioxo-5-(2-fluorophenyl)-3-isocyanate-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Phosgene in toluene (1.93M solution; 7ml) was added to a solution of the intermediate 3 (0.2g) in dichloromethane (3ml). The resulting solution was stirred at 23° for 5h, then concentrated in vacuo at 50° for 3h to give the title compound as a white solid (0.21g). M.p. 167-8°.

Intermediate 8

25 2,4-Dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)-3-(phenyloxycarbonylamino)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Pyridine (0.137ml) and phenyl chloroformate (0.21ml) were added to a solution of the intermediate 3 (0.3g) in dichloromethane (15ml) under a nitrogen atmosphere. The resulting solution was stirred at 23° for 30min, then washed with a 1% solution of hydrochloric acid (15ml), a 5% solution of sodium hydrogen carbonate (15ml) and brine

(20ml). The organic layer was dried and concentrated in vacuo to a solid which was triturated with ethyl acetate to give the title compound as a white solid (0.3g). M.p. 226-7°. T.l.c. CH-EA (1:1), Rf=0.75.

5 Intermediate 9

1-(3,3-Dimethylbut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-3-(phenyloxycarbonylamino)-
2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Pyridine (0.64ml) and phenyl chloroformate (1.0ml) were added to a solution of the intermediate 6 (1.5g) in dichloromethane (100ml) under a nitrogen atmosphere. The 10 resulting solution was stirred at 23° for 30min, then washed with a 1% solution of hydrochloric acid (2x70ml), a 5% solution of sodium hydrogen carbonate (2x70ml) and brine (100ml). The organic layer was dried and concentrated in vacuo to a solid which was triturated with diethyl ether to give the title compound as a white solid (1.4g). M.p. 199- 200°. T.l.c. CH-EA (1:1), Rf=0.82.

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Intermediate 10

2-(3,3-Dimethyl-2-hydroxybut-1-yl)amino-2'-fluorodiphenylamine

1,2-Epoxy-3,3-dimethylbutane (7ml) was added, portionwise, to a mixture of the 20 2-amino-2'-fluorodiphenylamine (7.46g) and p.toluenesulfonic acid (0.6g) in ethanol (30ml) heated to 80°. The mixture was stirred at 80° for 19h, then concentrated in vacuo and partitioned between water (100ml) and ethyl acetate (150ml). The organic layer was washed with a 5% solution of sodium hydrogen carbonate (2x100ml), brine (150ml), dried and concentrated in vacuo to an oil, which was purified by flash chromatography (eluting with CH-EA 80:20) to give the title compound as a yellow oil (3.21g). T.l.c. 25 CH-DCM (1:1), Rf 0.25.

Intermediate 11

1-(3,3-Dimethyl-2-hydroxybut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-3-phenylhydrazono-
2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 10 (1.8g) and the 2- phenylhydrazonomalonyldichloride (1.76g) were each taken up in THF (35ml) and dropped in a flask containing THF (30ml) maintained at - 15° under a nitrogen atmosphere. After complete addition the solution was allowed to warm to room temperature and then heated to 50° for 3h. The solution was concentrated in vacuo to an oil, which was purified by flash chromatography (eluting with CH-EA 8:2) to give the title compound as a yellow solid (2.1g). M.p.217-8°. T.l.c. CH-EA (2:1), Rf 0.71.

Intermediate 12

10 3-Amino-1-(3,3-dimethyl-2-hydroxybut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Zinc dust (2.17g) was added portionwise to a solution of the intermediate 11(2.1g) in glacial acetic acid (30ml) previously cooled to 0°. The mixture was stirred at 23° for 20h, then diluted with water (100ml) and decanted from zinc. Solid sodium carbonate was added until pH=9, and the mixture was extracted with ethyl acetate (3x100ml). The combined organic extracts were washed with brine (200ml), dried and concentrated in vacuo to an oil, which was purified by flash chromatography (eluting with EA) to give the title compound as a white foam (1.09g). M.p. 104-5° T.l.c. EA- methanol (20:2), Rf 0.66 and 0.61.

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Intermediate 13

3-Amino-1-(3,3-dimethyl-2-hydroxybut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine (diastereomer I; 13a and diastereomer II; 13b)

The diastereomeric mixture, intermediate 12 was separated by preparative HPLC (Column Spherisorb 5uCN 25x0.46cm) eluting with hexane/ethanol/isopropanol 85:10:5 and isopropylamine 0.05% (flux 2ml/min, detection UV at 235nm) to give the title compound 13a (retention time 8.9min) as a white solid (0.3g) M.p. 164-5° T.l.c. EA-methanol (20:2), Rf 0.66. and the title compound 13b (retention time 6min) as a white foam (0.35g). T.l.c. EA-methanol (20:2), Rf 0.61.

Intermediate 142-(1,3-dimethylbut-1-yl)amino-diphenylamine

Sodium borohydride (0.4g) was added portionwise to a mixture of 2-amino-diphenylamine (0.5g), sodium acetate trihydrate (0.5g) and 5 4-methyl-2-oxo-pentane (0.25ml) in acetic acid (1.7ml), water (5ml) and ethanol (4ml) cooled at 0°C. A further amount of sodium borohydride (2.0 g) and of 4-methyl-2-oxo-pentane (3ml) were added and the solution was stirred at 23° for 30 min., then diluted with ethyl acetate (100ml) and water (100ml). The organic layer was washed with a 10% solution of sodium hydroxide (50ml) and brine (50ml), dried and 10 concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 90:10) to give the title compound as a yellow oil (0.42g). T.l.c. CH-EA (90:10), Rf 0.79. IR :3420 (NH), 1599, 1514 and 1497 (C=C) cm-1;

Intermediate 151-(1,3-Dimethylbut-1-yl)-2,4-dioxo-5-phenyl-3-phenylhydrazono-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 14 (0.42g), and 2- phenylhydrazonomalonyldichloride (0.46g) were each taken up in THF (20ml) and dropped in a flask containing THF (10ml) maintained at 0° under a nitrogen atmosphere. After complete addition the solution 20 was allowed to warm to 23° and stirred for 20h. A further amount of 2-phenylhydrazonomalonyldichloride (0.13g) was added and stirring continued for 1 h at 23° and then at 50° for 90 min. The reaction mixture was diluted with ethyl acetate (200ml); the organic layer was washed with a 10% solution of sodium hydroxide (60ml) and brine (2x70 ml), dried and concentrated in vacuo to give an oil, which was purified 25 by flash chromatography (eluting with CH-EA 95:5, increasing polarity to 90:10) to give the title compound as a yellow solid (0.43g). T.l.c. CH-EA (70:30), Rf 0.73. IR :1668, 1653 (C=O); 1591 (C=C) cm-1;

Intermediate 16

3-Amino-1-(1,3-dimethylbut-1-yl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Zinc dust (0.55g) was added portionwise to a solution of the intermediate 15 (0.42g) in glacial acetic acid (10ml) cooled at 0°. The mixture was stirred at 23° for 8h, then

5 decanted from zinc, diluted with ethyl acetate (50 ml), washed with a 10% solution of sodium hydroxide (60 ml), brine (2x60ml), dried and concentrated in vacuo to give an oil which was purified by flash chromatography (eluting in gradient from CH-EA 1:1 to DCM-methanol 90:10) to give the title compound as a white foam (0.22g). T.l.c. DCM-methanol (90:10), Rf 0.53. IR :3500-3000 (NH2), 1703 and 1672 (C=O), 1593

10 (C=C) cm⁻¹;

Intermediate 17

2-Amino-5-chloro-diphenylamine

Potassium carbonate (29g) and sodium hydrosulphite (25.3g) were added portionwise

15 over 1 hour to a suspension of 5-chloro-2-nitrodiphenylamine (8g) in 95% ethanol (250ml) and water (250ml). The mixture was stirred at 23° for 20h, then a further amount of sodium hydrosulfite (1g) was added and stirring continued for 1h. The reaction mixture was acidified to pH4 with conc. hydrochloric acid and then a 10% solution of sodium hydroxide was added until the pH was 10. The solution was concentrated in vacuo and extracted with ethyl ether (2x250ml). The combined organic extracts were washed with brine (2x250ml), dried and concentrated in vacuo to give the crude compound as a yellow solid (7.8g) which was purified by flash chromatography (eluting with P-EE 1:1) to give the title compound as a yellow foam (4.4g). T.l.c. CH-EA (1:1), Rf 0.50. IR : 3412 and 3320 (NH), 1592-1589 (C=C) cm⁻¹;

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Intermediate 18

5-Chloro-2-(3-methylbut-1-yl)amino-diphenylamine

Sodium borohydride (2g) was added portionwise to a mixture of the intermediate 17 (2g), sodium acetate trihydrate (2.28g) and 3-methylbutyraldehyde (2ml) in acetic acid (8ml),

30 water (15ml) and ethanol (35ml) cooled to 0°C. The solution was stirred at 23° for 30

min, then diluted with ethyl acetate (200ml). The organic layer was washed with a 10% solution of potassium carbonate (100ml) and brine (100ml), dried and concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 95:5) to give the title compound as a yellow oil (0.8g). T.l.c. CH-EA (1:1), Rf 0.72.

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Intermediate 19

7-Chloro-2,4-dioxo-1-(3-methylbut-1-yl)-5-phenyl-3-phenylhydrazone-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 18 (1.15g) and 2-phenylhydrazonomalonyldichloride (1.17g) were each taken up in THF (30ml) and added dropwise in a flask containing THF (10ml) maintained at 0° under a nitrogen atmosphere. After complete addition the solution was allowed to warm to 23°C, stirred for 30 min., then heated at 60° for 2h. The solution was diluted with ethyl acetate (150ml), washed with brine (2x100ml), dried and concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 95:5 , increasing polarity to 70:30) to give the title compound as a yellow solid (1.12g). T.l.c. CH-EA (1:1), Rf 0.61. IR : 3452 (NH), 1664 (C=O) cm⁻¹;

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Intermediate 20

3-Amino-2,4-dioxo-7-chloro-5-phenyl-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

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A solution of the intermediate 19 (0.6g) in glacial acetic acid (14ml) was added, dropwise, to a suspension of zinc dust (0.76g) in glacial acetic acid (14 ml) cooled at 0°. The mixture was stirred at 23° for 3h, then decanted from zinc, washed with ethyl acetate (80 ml) and then with 10% sodium hydroxide (100ml) and brine (70ml).

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The combined organic extracts were dried and concentrated in vacuo to give an oil which was purified by flash chromatography (eluting in gradient from CH-EA 1:1 to EA -methanol 27:3) to give the title compound (0.3g). T.l.c. DCM-methanol (27:3), Rf 0.5.

Intermediate 21

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4-Chloro-2-nitrodiphenylamine

A mixture of 4-chloro-2-nitroaniline (5.5g), bromobenzene (20ml), potassium carbonate (1.63g) and copper(I) iodide (0.68g) was heated to 180° for 36h. The reaction mixture was cooled to room temperature, then ethyl acetate (200ml) and water (300ml) were added ; the organic extracts were washed with brine (2x150ml), dried and concentrated in vacuo to give the crude compound which was purified by flash chromatography (eluting with CH-EA 95:5) to give the title compound (3.67g) . T.l.c. CH-EA (1:1), Rf 0.71.

5 Intermediate 22

2-Amino-4-chlorodiphenylamine

10 Potassium carbonate (13g) and sodium hydrosulphite (11.4g) were added portionwise over 3 hour to a suspension of 4-chloro-2- nitrodiphenylamine (3.6g) in 95% ethanol (100ml) and water (100ml).The mixture was stirred at 23° for 20h. The reaction mixture was then acidified to pH=4 with conc. hydrochloric acid (20ml); then 10% solution of sodium hydroxide (80ml) was added until pH=10 and the solution extracted with ethyl acetate (2x150ml). The combined organic extracts were washed with brine (2x150ml), dried and concentrated in vacuo to give the crude compound as a yellow solid (7.8g) which was purified by flash chromatography (eluting with CH-EA 90:10 then 70:30) to give the title compound as a yellow foam (2.37g) . T.l.c. CH-EA (1:1), Rf 0.66.

15

20 Intermediate 23

4-Chloro-2-(3-methylbut-1-yl)amino-diphenylamine

Bromo 3-methylbutane (0.62ml) was added to a solution of the intermediate 22 (1.00g) and sodium iodide (0.7g) in dimethylformamide (40ml) under a nitrogen atmosphere. The solution was stirred at 120° for 12h, then cooled at 23°C, diluted with ethyl acetate (150ml) and washed with brine (3x100ml). The combined organic extracts were dried and concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 95:5) to give the title compound as a yellow oil (0.74g). T.l.c. CH-EA (1:1), Rf 0.76.

30 Intermediate 24

8-Chloro-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl-3-phenylhydrazone-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 23 (0.74g) and the 2- phenylhydrazonomalonyldichloride (0.75g) were each taken up in THF (15ml) and dropped in a flask containing THF (20ml) maintained at 0° under a nitrogen atmosphere. After complete addition, the solution was allowed to warm to 23°C, stirred for 30min., then heated at 60° for 2h. The solution was diluted with ethyl acetate (120ml), washed with brine (2x100ml), dried and concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 95:5 , increasing polarity to 70:30) to give the title compound as a yellow solid (0.91g).

10 T.I.c. CH-EA (1:1), Rf 0.68

Intermediate 25

3-Amino-8-chloro-2,4-dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

15 To the solution of intermediate 24 (0.9g) in glacial acetic acid (20ml) at 0°, zinc dust (1.14g) was added portionwise . The mixture was stirred at 23° for 1h, then decanted from zinc, washed with ethyl acetate (150ml) and then with 10% sodium hydroxide (150ml) and brine (100ml). The combined organic extracts were dried and concentrated in vacuo to an oil which was purified by flash chromatography (eluting in gradient from
20 CH-EA 1:1 to EA-methanol 27:3) to give the title compound (0.53g). T. I.c.
EA-methanol (27:3), Rf 0.6.

Intermediate 26

4,5-Dichloro-2-nitrodiphenylamine

25 A mixture of 4,5-dichloro-2-nitroaniline (5.0g), bromobenzene (16ml) potassium carbonate (1.17g) and copper(I) iodide (0.46g) was heated to 150° for 36h. The reaction mixture was concentrated in vacuo to give the crude compound which was purified by flash chromatography (eluting with CH-EA 90:10) to give the title compound (4.34g)
T.I.c. CH-EA (1:1), Rf 0.7.

Intermediate 272-Amino4,5-dichloro-diphenylamine

Potassium carbonate (13.8g) and sodium hydrosulfite (12.1g) were added portionwise over 3 hour to a suspension of 4,5-dichloro-2-nitrodiphenylamine (4.34g) in 95% ethanol (100ml) and water (100ml). The mixture was stirred at 23° for 20h. The reaction mixture was then acidified to pH=4 with conc. hydrochloric acid (20ml), then 10% solution of sodium hydroxide (80 ml) was added until pH=10 and the solution extracted with ethyl acetate (2x120ml). The combined organic extracts were washed with brine (2x100ml), dried and concentrated in vacuo to give the crude compound which was purified by flash chromatography (eluting with CH-EA 90:10 then 80:20) to give the title compound as a yellow foam (2.15g). T.l.c. CH-EA (1:1), Rf 0.54.

Intermediate 284,5-Dichloro-2-(3-methylbut-1-yl)amino-diphenylamine

1-Bromo-3-methylbutane (1.2ml) was added to a solution of the intermediate 27 (2.15g) and sodium iodide (1.3g) in dimethylformamide (70ml) under a nitrogen atmosphere. The solution was stirred at 120° for 9h, and at 23°C for 20h. A further amount of bromo-3-methylbutane (0.5ml) was then added and stirring was continued at 120° for 8h. The reaction mixture was diluted with ethyl acetate (300ml) and washed with brine (150ml). The combined organic extracts were dried and concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 95:5) to give the title compound as a yellow oil (1.72g). T.l.c. CH-EA (1:1), Rf 0.70.

Intermediate 297,8-dichloro-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl-3-phenylhydrazone-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 28 (1.72g) and the 2-phenylhydrazonomalonyldichloride (1.53g) were each taken up in THF (15ml) and dropped in a flask containing THF (40ml) maintained at 0° under a nitrogen atmosphere. After complete addition the solution was allowed to warm at 23°C, stirred for 45 min., then heated at 60° for 1h and 30min. The solution was

diluted with ethyl acetate (150ml), washed with brine (2x100ml), dried and concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 95:5, increasing polarity to 80:20) to give the title compound as a yellow solid (1.85g).
T.l.c. CH-EA (1:1), Rf 0.66.

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Intermediate 303-Amino-7-8-dichloro-2,4-dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

To the solution of the intermediate 29 (1.0g) in glacial acetic acid (15ml) at 0°, zinc dust (0.65g) was added portionwise. The mixture was stirred at 23° for 6h, then decanted from zinc, washed with ethyl acetate (150 ml) and then with 10% sodium hydroxide (150ml) and brine (100ml). The combined organic extracts were dried and concentrated in vacuo to give an oil which was purified by flash chromatography (eluting in gradient from CH-EA 1:1 to EA-methanol 80:20) to give the title compound (0.44g). T.l.c.
EA-methanol (27:3), Rf 0.59.

Intermediate 314-Fluoro-2-nitrodiphenylamine

A mixture of 4-Fluoro-2-nitroaniline (5.0g), bromobenzene (20ml), potassium carbonate (1.54g) and copper(I) iodide (0.61g) was heated to 150° for 30h. The reaction mixture was cooled at 23°C, then ethyl acetate (200 ml) was added; the organic extracts were washed with brine (100ml), dried and evaporated in vacuo to give the crude compound which was purified by flash chromatography (eluting with CH-EA 95:5) to give the title compound (2.4g). T.l.c. CH-EA (1:1), Rf 0.68.

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Intermediate 322-amino-4-Fluoro-diphenylamine

Potassium carbonate (9.3g) and sodium hydrosulfite (8.2g) were added portionwise over 3 hour to a suspension of 4-fluoro-2- nitrodiphenylamine (2.4g) in 95% ethanol (70ml) and water (70ml). The mixture was stirred at 23° for 20h. The reaction mixture was the n

acidified to pH=4 with conc. hydrochloric acid (15ml), then 10% solution of sodium hydroxide (50 ml) was added until pH=10, and the concentrated solution extracted with ethyl acetate (2x100ml). The combined organic extracts were washed with brine (2 x80ml), dried and concentrated in vacuo to give the crude compound which was purified by flash chromatography (eluting with CH-EA 90:10 then 80:20) to give the title compound as a yellow foam (1.44g) . T.l.c. CH-EA (1:1), Rf 0.72.

Intermediate 33

4-Fluoro-2-(3-methylbut-1-yl)amino-diphenylamine

10 1-Bromo 3-methylbutane (1.0ml) was added to a solution of the intermediate 32 (1.44g) and sodium iodide (1.1g) in dimethylformamide (60ml) under a nitrogen atmosphere. The solution was stirred at 120° for 9h; the reaction mixture was diluted with ethyl acetate (300ml) and washed with brine (3x150ml). The combined organic extracts were dried and concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 95:5) to give the title compound as a yellow oil (0.96g). T.l.c. CH-EA (1:1), Rf 0.74.

Intermediate 34

2,4-Dioxo-8-fluoro-1-(3-methylbut-1-yl)-5-phenyl-3-phenylhydrazone-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

20 The intermediate 33 (0.96g) and 2-phenylhydrazonomalononyldichloride (1.01g) were each taken up in THF (15ml) and dropped in a flask containing THF (40ml) maintained at -0° under a nitrogen atmosphere. After complete addition the solution was allowed to warm at 23°C, stirred for 30min., then heated at 60° for 2h. The solution was diluted with ethyl acetate (120ml), washed with brine (2x100ml), dried and concentrated in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 95: 5 , increasing polarity to 80:20) to give the title compound as a yellow solid (1.3g). T.l.c. CH-EA (1:1), Rf 0.74.

Intermediate 353-Amino-2,4-dioxo-8-Fluoro1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

To the solution of the intermediate 34 (1.3g) in glacial acetic acid (20ml) at 0°, zinc dust (1.2g) was added portionwise. The mixture was stirred at 23° for 1h, then decanted from zinc, washed with ethyl acetate (150ml) and then with 10% sodium hydroxide (150ml) and brine (100ml). The combined organic extracts were dried and concentrated in vacuo to give an oil which was purified by flash chromatography (eluting in gradient from CH-EA 1:1 to EA-methanol 80:20) to give the title compound (0.72 g). T.l.c.

EA-methanol (27:3), R_f 0.47.

Intermediate 362,4-Dioxo-5-phenyl-1-(2-phenylethyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Sodium hydride 80% dispersion in oil (0.13g) was added portionwise to a solution of the 15 2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine (compound a) (1g) in DMF (18ml) previously cooled at 0°. The reaction was stirred for 20min at 0°, then a solution of 2-phenylethyl bromide (0.85ml) in DMF (2ml) was added dropwise, the mixture was stirred at 23° for 15h, then diluted with EA (80ml) and washed with brine (3x100ml), dried and concentrated "in vacuo". The crude product was purified by flash 20 chromatography (eluting with CH-EA 1:1) to give the title compound as a white powder (0.85g). T.l.c. CH-EA (1:1), R_f=0.27.

Intermediate 373-Azido-2,4-dioxo-5-phenyl-1-(2-phenylethyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 36 (0.85g) in THF (20ml), cooled to -70°, was added dropwise to a solution of potassium tert-butoxide (0.3g) in THF (10ml) cooled to -70°, under a nitrogen atmosphere. The mixture was stirred for 20min at -70°, then a solution of 2,4,6-trisopropylbenzenesulphonyl azide (0.96g) in THF (15ml), previously cooled to -70° and acetic acid (0.14ml) were added. The reaction mixture was allowed to stand at

23° and stirred for 1.5 h, then more acetic acid (0.14ml) was added and the mixture was stirred for 2 h. Ethyl acetate (150ml) was added and the solution was washed with a saturated solution of sodium hydrogen carbonate (100ml) and brine (3x100ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 90:10) to give the title compound as a white foam (0.38g). T.l.c. CH-EA (1:1), $R_f=0.57$.

Intermediate 38

3-Amino-2,4-dioxo-5-phenyl-1-(2-phenylethyl)-2,3,4,5-tetrahydro-1H-1,5-

10 benzodiazepine

A solution of the intermediate 37 (0.38g) in ethanol (15ml) and ethyl acetate (15ml) was stirred under hydrogen, at 1 atm., in presence of 5% Pd/CaCO₃ (0.25g), at 23°, for 3h. The catalyst was filtered off on a pad of celite, washing with dichloromethane (25ml) and ethanol (25ml) and the organic layer was concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with DCM-ethanol 90:10) to give the title compound as a white foam (0.3g). T.l.c. DCM-ethanol (90:10), $R_f=0.1$.

Intermediate 39

1-(1-Adamantyl)methyl-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

20 Sodium hydride 80% dispersion in oil (0.07g) was added portionwise to a solution of the compound (a) (0.5g) in DMF (50ml). The reaction mixture was stirred for 30min, then a solution of 1-adamantylmethyl methanesulfonate (0.537g) in DMF (3ml) was added. The reaction mixture was stirred at 120° for 7h and at 23° for 15h, then concentrated. The residue was diluted with ethyl acetate (100ml) washed with brine (2x30ml) and water (50ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 1:1) to give the title compound as a white foam (0.15g). T.l.c. CH-EA (1:1), $R_f=0.42$.

Intermediate 40

1-(1-Adamantyl)methyl-1-(2-phenylethyl)-3-azido-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of potassium tert-butoxide (0.146g) in THF (7ml) was added dropwise to a solution of the intermediate 39 (0.4g) in THF (15ml), cooled to -70°, under a nitrogen atmosphere. The mixture was stirred for 20 min at -70°, then a solution of 2,4,6-triisopropylbenzenesulphonyl azide (0.53g) in THF (7ml), previously cooled to -70° and acetic acid (0.14ml) were added. The reaction mixture was allowed to stand at 23° and stirred for 15h, ethyl acetate (70ml) was then added and the solution was washed with water (2x50ml) and brine (2x30ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 70:30) to give the title compound as a white foam (0.338g). T.l.c. CH-EA (1:1), R_f=0.73.

Intermediate 41

1-(1-Adamantyl)methyl-3-amino-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 40 (0.18g) in ethanol (10ml) and ethyl acetate (5ml) was stirred under hydrogen, at 1 atm., in presence of 5% Pd/CaCO₃ (0.2g), at 23°, for 3h, then the catalyst was filtered off on a pad of celite and the organic layer was concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with DCM-methanol 90:10) to give the title compound as a white foam (0.15g). T.l.c. DCM-methanol (90:10), R_f=0.51.

Intermediate 42

1-(2,2-Dimethylethoxycarbonylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Sodium hydride 80% dispersion in oil (0.155g) was added portionwise to a solution of the compound (a) (1.022g) in DMF (30ml) previously cooled to 0°. The reaction was stirred for 15 min at 23°, then t- butyl bromoacetate (0.7ml) was added. The solution was stirred at 23° for 1h, then brine (100ml) was added and the mixture extracted with ethyl acetate (3x30ml), dried and concentrated "in vacuo". The crude product was purified by flash

chromatography (eluting with CH-EA 60:40) to give the title compound as a white powder (1.31g). T.l.c. CH-EA (60:40), $R_f=0.4$.

Intermediate 43

5 3-Azido-1-(2,2-dimethylethoxycarbonylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 42 (0.5g) in THF (6ml), cooled to -70°, was added dropwise to a solution of potassium tert-butoxide (0.168g) in THF (6ml) cooled to -70°, under a nitrogen atmosphere. The reaction mixture was stirred for 30min at -70°, then a 10 solution of 2,4,6-triisopropylbenzenesulphonyl azide (0.556g) in THF (6ml), previously cooled to -70° and acetic acid (0.078ml) were added. The reaction mixture was allowed to stand at 23° and stirred for 18h, ethyl acetate (30ml) was added and the solution was washed with brine (3x100ml), a saturated solution of sodium hydrogen carbonate (20ml), brine (20ml), dried and concentrated "in vacuo". The crude product was purified by flash 15 chromatography (eluting with CH-EA 70:30) to give the title compound as a white foam (0.5g). T.l.c. CH-EA (1:1), $R_f=0.36$.

Intermediate 44

20 3-Amino-1-(2,2-dimethylethoxycarbonylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 43 (0.354g) in a mixture of ethanol (10ml) and ethyl acetate (2ml) was stirred under hydrogen, at 1 atm., in presence of 5% Pd/CaCO₃ (0.183g), at 23°, for 3h, then more 5% Pd/CaCO₃ (0.183g) was added and the reaction stirred for 15h. The catalyst was filtered off on a pad of celite, washing with 25 dichloromethane (9ml) and methanol (5ml) and the organic layer was concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with DCM-methanol 96:4) to give the title compound as a white foam (0.33g). T.l.c. DCM-methanol (95:5), $R_f=0.5$.

30 Intermediate 45

1-(3,3-Dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Sodium hydride 80% dispersion in oil (0.100g) was added portionwise to a solution of the compound (a) (0.7g) in DMF (60ml). The reaction mixture was stirred for 30min, then a solution of 3,3-dimethylbutyl methanesulfonate (0.575g) in DMF (3ml) was added. The reaction mixture was stirred at 90° for 50 min, at 23° for 15h, at 90° for 2h and at 140° for 45min, then concentrated. The residue was diluted with water (30ml) and brine (20ml) and extracted with ethyl acetate (150ml); the organic layer was washed with water (2x50ml) and brine (50ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 1:1) to give the title compound as a white foam (0.4g). T.l.c. CH-EA (1:1), R_f=0.39.

Intermediate 463-Azido-1-(3,3-dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of potassium tert-butoxide (0.146g) in THF (7ml), cooled to -70°, was added dropwise to a solution of the intermediate 45 (0.4g) in THF (15ml), cooled to -70°, under a nitrogen atmosphere. The solution was stirred for 20 min at -70°, then a solution of 2,4,6-triisopropylbenzenesulphonyl azide (0.530g) in THF (7ml), previously cooled to -70° and acetic acid (0.139ml) were added. The reaction mixture was allowed to stand at 23° and stirred for 18h, then ethyl acetate (75ml) was added and the solution was washed with water (2x50ml) and brine (2x30ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 30:70) to give the title compound as a white foam (0.338g). T.l.c. CH-EA (1:1), R_f=0.73.

25 Intermediate 473-Amino-1-(3,3-dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 46 (0.298g) in a mixture of ethanol (18ml) and ethyl acetate (7ml) was stirred under hydrogen, at 1 atm., in presence of 5% Pd/CaCO₃, (0.186g), at 23°, for 1.5 h, then more 5% Pd/CaCO₃, (0.180g) was added and the reaction

stirred for 1h. The catalyst was filtered off on a pad of celite, washing with ethanol (20ml) and the organic layer was concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with DCM-methanol 90:10) to give the title compound as a white foam (0.205g). T.l.c. DCM-methanol (90:10), $R_f=0.46$.

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Intermediate 481-(3,3-Dimethylbutyl)-2,4-dioxo-3-isocyanato-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

To a solution of the intermediate 47 (0.3g) in dichloromethane (20ml), a 1.93M solution of COCl₂ in toluene (10ml) was added. The reaction mixture was stirred for 5h at 23°, then concentrated "in vacuo" at 50° for 3h to obtain the title compound as a white foam (0.370g). IR: 2218 (N=C=O); 1693, 1668 (C=O), (C=C) cm⁻¹;

Intermediate 491-(3,3-Dimethylbutyl)-2,4-dioxo-5-phenyl-3-phenyloxycarbonylamino-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

To a solution of the intermediate 47 (1g) in dichloromethane (50ml), pyridine (0.46ml) and phenylchloroformate (0.7ml) were added. The reaction mixture was stirred for 30min at 23°, then washed with a 1% solution of hydrochloric acid (20ml), a 5% solution of sodium hydrogen carbonate (20ml), water (20ml), brine (20ml), dried and concentrated "in vacuo". The crude product was triturated with acetonitrile (10ml) to obtain the title compound as a white powder (1.2g). T.l.c. CH-EA (1:1) $R_f=0.8$.

Intermediate 501-[2-(1-adamantyl)ethyl]-2,4-dioxo-5-phenyl-3-phenyloxycarbonylamino-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

To a solution of the intermediate 53 (0.1g) in dichloromethane (10ml), pyridine (0.03ml) and phenylchloroformate (0.01ml) were added. The reaction mixture was stirred for 2h at 23°, then diluted with dichloromethane (30ml), washed with a saturated solution of ammonium chloride (30ml) and brine (40ml), dried and concentrated "in vacuo". The

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crude product was triturated with acetonitrile (10ml) to obtain the title compound as a white powder (0.05g). T.l.c. CH-EA (1:1) $R_f=0.77$.

Intermediate 51

5 1-[2-(1-Adamantyl)ethyl]-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine
Sodium hydride 80% dispersion in oil (0.15g) was added portionwise to a solution of the compound (a) (0.8g) in DMF (20ml) previously cooled to 0°. The reaction was stirred for 15min at 0°, a solution of 2-(1-adamantyl)ethyl bromide (0.8g) in DMF (10ml) was added dropwise, the mixture was stirred at 23° for 8h, then diluted with DMF (20ml) heated at
10 80° for 1h and allowed to stand at 23° for 2 days. ethyl acetate (200ml) was added and the solution was washed with brine (3x100ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 70:30) to give the title compound as a white foam (0.45g). T.l.c. CH-EA (1:1), $R_f=0.42$.

15 Intermediate 52

1-[2-(1-Adamantyl)ethyl]-3-azido-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of potassium tert-butoxide (0.2g) in THF (10ml) was added dropwise to a solution of the intermediate 51 (0.67g) in THF (20ml), cooled to -70°, under a nitrogen atmosphere. The mixture was stirred for 20 min at -70°, then a solution of 2,4,6-triisopropylbenzenesulphonyl azide (0.65g) in THF (10ml), previously cooled to -70° and acetic acid (0.18ml) were added. The reaction mixture was allowed to stand at 23° and stirred for 15h, then EA (150ml) was added and the solution was washed with a 5% solution of sodium hydrogen carbonate (80ml) and brine (100ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 90:10) to give the title compound as a white foam (0.71g). T.l.c. CH-EA (1:1), $R_f=0.68$.

Intermediate 53

1-[2-(1-Adamantyl)ethyl]-3-amino-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 52 (0.71g) in ethanol (30ml) and ethyl acetate (15ml) was stirred under hydrogen, at 1 atm., in presence of 5% Pd/CaCO₃ (0.7g), at 23°, for 3h. The catalyst was filtered off on a pad of celite, washing with methanol (50ml) and the organic layer was concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with DCM-methanol 90:10) to give the title compound as a white foam (0.5g). T.l.c. DCM- methanol (90:10), R_f=0.62.

10 Intermediate 54

1-(2,3-Dimethyl)butyl-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5- benzodiazepine

Sodium hydride 80% dispersion in oil (0.06g) was added portionwise to a solution of the compound (a) (0.38g) in DMF (10ml). The reaction was stirred at 23° for 1h, then 2,3-dimethylbutyl methanesulfonate (0.32g) was added. The mixture was stirred at 23° for 15h, then water (70ml) was added and the solution was extracted with ethyl acetate (2x50ml) and the combined organic layer were washed with brine (2x50ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH- EA 80:20) to give the title compound as a white foam (0.23g). T.l.c. CH-EA (1:1), R_f=0.4.

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Intermediate 55

3-Azido-1-(2,3-dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of potassium tert-butoxide (0.121g) in THF (10ml) was added to a solution of the intermediate 54 (0.33g) in THF (20ml), cooled to -70°, under a nitrogen atmosphere. The mixture was stirred for 30 min at -70°, then a solution of 2,4,6-triisopropylbenzenesulphonyl azide (0.349g) in THF (10ml), previously cooled to -70° and, after 20 min, acetic acid (0.06ml) were added. The reaction mixture was allowed to stand at 23° and stirred for 24h; ethyl acetate (50ml) was added and the solution washed with a 5% solution of sodium hydrogen carbonate (2x50ml) and brine (2x50ml), dried and

concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 80:20) to give the title compound as a white foam (0.1g). T.l.c. CH-EA (1:1), $R_f=0.53$.

5 Intermediate 56

3-Amino-1-(2,3-dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 55 (0.19g) in ethanol (15ml) and ethyl acetate (3ml) was stirred under hydrogen, at 1 atm., in presence of 5% Pd/CaCO₃, (0.18g), at 23°, for 4h.

10 The catalyst was filtered off on a pad of celite, washed with ethyl acetate and the organic layer was concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with EA-methanol 90:10) to give the title compound as a white foam (0.95g). T.l.c. EA-methanol (90:10), $R_f=0.55$.

15 Intermediate 57

1-Butyl-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Sodium hydride 80% dispersion in oil (0.031g) was added portionwise to a solution of the compound (a) (0.3g) in DMF (18ml) at 0°, under a nitrogen atmosphere. The reaction was allowed to stand at 23° for 30 min, then a solution of 1-bromobutane (0.154ml) in

20 DMF (3ml) was added dropwise. The mixture was stirred at 23° for 2h, then water (30ml) was added and the solution was extracted with ethyl acetate (2x60ml) and the combined organic layer were dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 60:40) to give the title compound as a white foam (0.057g). T.l.c. CH-EA (70:30), $R_f=0.53$.

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Intermediate 58

3-Azido-1-butyl-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of potassium tert-butoxide (0.0418g) in THF (2ml) was added to a solution of the intermediate 57 (0.104g) in THF (2ml), cooled to -70°, under a nitrogen atmosphere.

30 The mixture was stirred for 30 min at -70°, then a solution of 2,4,6-

triisopropylbenzenesulphonyl azide (0.136g) in THF (10ml), previously cooled to -70° and acetic acid (0.019ml) were added. The reaction mixture was allowed to stand at 23° and stirred for 24 hrs, ethyl acetate (50ml) added and the solution washed with a 5% solution of sodium hydrogen carbonate (10ml), brine (10ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 80:20) to give the title compound as a white foam (0.043g). T.l.c. CH-EA (60:40), R_f=0.67.

Intermediate 59

10 3-Amino-1-butyl-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine
A solution of the intermediate 58 (0.217g) in ethanol (4ml) and ethyl acetate (10ml) was stirred under hydrogen, at 1 atm., in presence of 5% Pd/CaCO₃, (0.18g), at 23°, for 10h. The catalyst was filtered off on a pad of celite, washed with ethyl acetate (3x5ml) and ethanol (3x5ml) and the organic layer was concentrated "in vacuo". The crude product
15 was purified by flash chromatography (eluting with EA-methanol 90:10) to give the title compound as a white foam (0.094g). T.l.c. EA-methanol (95:5), R_f=0.25.

Intermediate 60

20 2,4-Dioxo-5-phenyl-1-(3-methyl-2-oxo)butyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine
Sodium hydride 80% dispersion in oil (0.4g) was added portionwise to a solution of the compound (a) (2g) in DMF (50ml), previously cooled to 0 C. The reaction was stirred for 15 min at 0°, then a solution of 1-bromo-3-methyl-2-oxobutane (2.6g) in DMF (10ml) was added dropwise, the mixture was stirred at 0 C for 45 min, ethyl acetate (450ml) added and the solution washed with brine (4x100ml), dried and concentrated "in vacuo".
25 The crude product was purified by flash chromatography (eluting with CH-EA 1:1) to give the title compound as a white foam (2.3g). T.l.c. CH-EA (1:1), R_f=0.19.

3

Intermediate 61

30 3-Azido-2,4-dioxo-1-(3-methyl-2-oxo)butyl-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of potassium tert-butoxide (0.185g) in THF (10ml) was added dropwise to a solution of the intermediate 60 (0.5g) in THF (20ml), cooled to -70°, under a nitrogen atmosphere. The mixture was stirred for 20 min at -70°, then a solution of 2,4,6-triisopropylbenzenesulphonyl azide (0.688g) in THF (10ml), previously cooled to -70° and acetic acid (0.2ml) were added. The reaction mixture was allowed to stand at 23° and stirred for 15h, ethyl acetate (400ml) added and the solution was washed with brine (3x100ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 1:1) to give the title compound as a foam. T.l.c. CH-EA (1:1), R_f=0.51.

10

Intermediate 62

3-Amino-2,4-dioxo-1-(3-methyl-2-oxo)butyl-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 61 (0.85g) in ethanol (35ml) was stirred under hydrogen, at 1 atm., in presence of 5% Pd/CaCO₃, (1g), at 23°, for 2h. The catalyst was filtered off on a pad of celite, washing with ethanol (30ml) and the organic layer was concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with DCM-methanol 90:10) to give the title compound as a white foam (0.5g). T.l.c. DCM-ethanol (95:5), R_f=0.56.

20

Intermediate 63

N-[2,4-Dioxo-1-(3-methyl-2-oxo)butyl-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.2ml) was added to a solution of the intermediate 62 (0.43g) in dry acetonitrile (15ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h and the formed precipitate filtered washing with acetonitrile (30ml) to give the title compound as a white solid (0.37g). T.l.c. CH-EA (1:1), R_f=0.27.

Intermediate 64

2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

30

NaH 80% dispersion in oil (0.057g) was added to a solution of 2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine (0.40g) in dry DMF (15ml). The reaction mixture was cooled at 0° and stirred for 15 min, 1-bromo-3-methyl-butane (0.23ml) in dry DMF (4ml) was added and stirring continued for 2h. The reaction mixture 5 was then diluted with water (100ml), extracted with ethyl acetate (3x100ml), washed with brine (2x50ml), dried and concentrated in vacuo to give an oil (0.75g) which was purified by flash chromatography (eluting with CH-EA 60:40) to give the title compound as a white solid (0.44g). T.l.c. CH-EA (1:1), Rf. 0.36

10 Intermediate 65

3-Azido-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A solution of the intermediate 64 (0.397g) in dry THF (7ml) was added to potassium tert-butoxide (0.154g) in dry THF (6ml) cooled at -78°. The reaction mixture was stirred 15 for 30min, then a cooled (- 78°) solution of 2,4,6 -triisopropyl-benzenesulphonylazide (0.49g) in dry THF (7ml) was added . After 5 min glacial acetic acid (0.07ml) was added and the solution was allowed to warm at 23° and stirred for 24 h. The reaction mixture was diluted with ethyl acetate (40ml) and washed with water (20ml) saturated sodium hydrogen carbonate solution (20ml) and brine (20ml). The combined organic extracts 20 were dried and concentrated in vacuo to give an oil (0.7g). Purification by flash chromatography (eluting with CH-EA 60:40) gave the title compound as a white solid (0.25g) T.l.c. CH- EA (60;40), Rf 0.3.

Intermediate 66

25 3-Amino-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

5% Pd/CaCO₃ (0.61g) was added to a solution of intermediate 65 (1.21g) in ethyl acetate (60ml) and ethanol (60ml) and the reaction mixture was hydrogenated at 1 atm for 3h and 30 min. The catalyst was filtered off and the solvent evaporated in vacuo to give the title 30 compound as a pale yellow foam (1.14g). T.l.c. DCM-methanol (95:5) , Rf 0.55.

Intermediate 67

3-Amino-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl--2,3,4,5-tetrahydro-1H-1,5-benzodiazepine (1S)-(+) -10-camphorsulphonic salt

5 To intermediate 66 (2.05g) dissolved in hot ethyl acetate (35 ml),
(1S)-(+) -10-camphorsulphonic acid was added. The resulting salt (5b) was crystallized
out from the cool solution by dropwise addition of cyclohexane; the precipitate was
filtered off and washed with cold cyclohexane to give a (+)/(-) 3/97 mixture of
diastereomeric salt (1.11g) and mother liquors. Recrystallization (twice) from
10 2-propanol afforded the pure title compound (0.49g). IR : 2750-2600 (NH3); 1736
, 1713, 1700 (C=O) cm-1; ¹H-NMR : 9.0-7.4 (m); 7.5 (d): 7.45-7.2 (m); 7.18 (t); 6.97
(d); 5.05 (s); 4.58 (m); 3.68 (m); 3.20 (m); 2.72 (m); 2.42 (m); 2.22 (m); 2.0 (m); 1.2 (m);
1.0-0.7(m).

15 Intermediate 68

(-)3-Amino-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl--2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Intermediate 67 (0.47g) was suspended in ethyl acetate and washed with 5% ammonia
solution (2x20 ml) and brine (2x20 ml). The organic layer was dried and concentrated in
20 vacuo to give the title compound as a white foam (0.27g). T.l.c. DCM-methanol (9 5:5) ,
Rf 0.55. [alpha]_D = -114. IR : 3377 (NH2), 1705- 1670 (C=C), 1593 cm-1; ¹H-NMR
: 7.5- 7.1(m); 6.95 (dd); 4.55(m), 4.23 (s); 3.7 (m); 1.8(m); 1.64- 1.4(m) ; 0.92(d); 0.89
(d) .

25 Intermediate 69

3-Amino-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl--2,3,4,5-tetrahydro-1H-1,5-benzodiazepine (1R)-(-) -10-camphorsulphonic salt

The mother liquors obtained after the initial precipitation of intermediate 67 were
evaporated to dryness to give a solid (2.19g). The residue was taken up in ethyl acetate
30 (30ml), extracted with a 5% ammonia solution (20ml) and washed with brine (20ml), the

organic layer dried and evaporated in vacuo to give a residue (1.0g).

(1R)-(-)-10-camphorsulphonic acid in ethyl acetate (6ml) was added to the solution of the residue (1g) in ethyl acetate (5ml) and the resulting solution was stirred at 0° for 2h. The obtained precipitate was filtered off, washed with ethyl acetate (20ml) and dried to give the title compound (0.97g). 1H-NMR : 9.0-7.2(m); 7.5 (d); 7.45-7.2 (m); 7.18 (t); 6.97 (d); 5.04 (s); 4.6 (m); 3.68 (m); 3.20 (m); 2.70 (m); 2.42 (m); 2.22 (m); 2.0 -1.8(m); 1.7-1.2 (m); 1.0-0.7(m).

Intermediate 70

10 (+)-3-Amino-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Intermediate 69 (0.95g) was suspended in ethyl acetate (130 ml), washed with a 5% ammonia solution (70ml) and stirred at 23° for 10 min. The organic layer was separated, washed with brine (3x70ml) dried and concentrated in vacuo to give the crude compound. Purification by flash chromatography (eluting with acetone-methanol 9:1) gave the title compound as a white foam (0.51g). [alpha]_D; IR : 3375 (NH₂), 1715-1661(C=C), 1591 cm-1; 1H-NMR : 7.5- 7.1(m); 6.95 (dd); 4.6-4.5(m), 4.24 (s); 3.8-3.65(m); 1.8(m); 1.62-1.4(m); 0.92(d); 0.89 (d).

20 Intermediate 71

3-Amino-2,4-dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine, (1R)-(-)-10-camphorsulfonate

A hot solution of (1R)-(-)-10-camphorsulfonic acid (1.685g) in ethyl acetate (15ml) was added, dropwise over 30', to a solution of the intermediate 3 (3.0g) in ethyl acetate (7ml) previously heated to 90 to under a nitrogen atmosphere. The resulting solution was heated to 90 for 10', then concentrated in vacuo. The residue triturated with EE-petroleum gave a (+)/(-) 50/50 mixture of diastereomeric salt (4.65g).

Recrystallization from 2-propanol gave the title compound (0.9g). M.p.216-7.
[alpha]D=+67.8.

Intermediate 723-(+)-Amino-2,4-dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Intermediate 70 (0.85g) was dissolved in a 5% ammonia solution (50ml) and extracted
5 with ethyl acetate (2x40ml). The combined organic extracts were washed with brine (60ml), dried and concentrated in vacuo to give the title compound as a white foam (0.5g). M.p. 125-6°.

T.l.c. DCM-methanol (30:1), Rf 0.38. [alpha]D=+115.2.

10 Intermediate 732-(Adamant-2-yl)amino-diphenylamine

Sodium borohydride (1.873g) was added portionwise to a mixture of 2-aminodiphenylamine 1 (0.61g), sodium acetate trihydrate (1.36g) and 2-adamantanone (0.5g) in acetic acid (2.1ml), water (8ml) and ethanol (6.5ml) cooled to 0. The reaction
15 mixture was stirred at 23 for 1h, then diluted with ethyl acetate (100ml). The organic layer was washed with water (30ml), a 10% solution of sodium hydroxide (2x25ml), water (30ml) and brine (20ml), dried and concentrated in vacuo to yield a residue which was taken up in DCM and the unreacted solid 2-adamantanone was removed by filtration.
The filtrate was concentrated in vacuo and purified by flash chromatography (eluting with
20 CH-EA 95:5) to give the title compound as a yellow solid (0.185g).

T.l.c. CH-EA (90:10), Rf 0.73.

Intermediate 741-(Adamant-2-yl)-2,4-dioxo-5-phenyl-3-phenylhydrazono-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 73 (0.96g) and 2-phenylhydrazonomalonyldichloride (0.89g) were each taken up in THF (10ml) and dropped in a flask containing THF (50ml) maintained at 0 under a nitrogen atmosphere. After complete addition the solution was allowed to warm to room temperature and then heated to 50 for 3h. The reaction mixture was concentrated

in vacuo to give an oil, which was purified by flash chromatography (eluting with CH-EA 90:10) to give the title compound as a yellow solid (0.758g).
T.l.c. CH-EA (80:20), Rf 0.60.

5 Intermediate 75

1-(Adamant-2-yl)-3-amino-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A suspension of the intermediate 74 (0.745g) in glacial acetic acid (10ml) was added to a mixture of zinc dust (0.956) in glacial acetic acid (5ml), cooled to 0. The mixture was 10 stirred at 23 for 3h, then diluted with water (100ml) and decanted from zinc. Solid sodium carbonate was added until pH=9 and the mixture extracted with ethyl acetate (3x100ml). The combined organic extracts were dried and concentrated in vacuo. The residue was triturated with ethyl acetate to give the title compound as a white solid (0.51g). M.p. 231-3°(dec). T.l.c. DCM-methanol (90:10), Rf 0.61.

15

Intermediate 76

1-(Adamant-2-yl)-2,4-dioxo-3-isocyanate-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Phosgene in toluene (1.93M solution, 10ml) was added to a solution of the intermediate 20 75 (0.285g) in dichloromethane (10ml); the resulting solution was stirred at 23 for 4h, then concentrated in vacuo at 50 for 2.5h to give the title compound as a white foam (0.29g).

IR : 2220 (N=C), 1697 and 1676 (C=O) cm⁻¹;

¹H-NMR : 7.50-7.15 (m); 7.05-6.95(m); 4.7 (s); 4.55 (m); 3.05 (m); 2.35 (m); 1.95-1.1 (m).

Intermediate 77

2-(2-cyclopentyl-ethyl)-amino-2'-fluoro-diphenylamine

Sodium borohydride (17.86g) was added portionwise to a mixture of 30 2-amino-2'-fluoro-diphenylamine (6.47g), sodium acetate trihydrate (4.24g) and

cyclopentylacetaldehyde (3.58g) in acetic acid (19.6ml), water (76ml) and ethanol (60ml) cooled to 0°. The reaction mixture was stirred at 23° for 1h and 30 min., then diluted with ethyl acetate (200ml). The organic layer was washed with water (70ml), a 10% solution of sodium hydroxide (70ml), and brine (50ml), dried and concentrated in vacuo
5 to yield a residue which was purified by flash chromatography (eluting with CH-EA 9:1) to give the title compound as a yellow oil (3.35g). T.l.c. CH-EA (9:1), Rf 0.78

Intermediate 78

10 1-(2-cyclopentyl-ethyl)-2,4-dioxo-5-(2-fluorophenyl)-3-phenylhydrazono-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 77 (3.30g) and 2-phenylhydrazonomalonyl dichloride (3.25g) were each taken up in THF (25ml) and dropped in a flask containing THF (150ml) maintained at 0° under a nitrogen atmosphere. After complete addition the solution was allowed to warm to 23°C; the reaction mixture was then heated to 55° for 3h and concentrated in vacuo.
15 The residue was taken up in cyclohexane/EA 7/3 (40 ml); the precipitate was filtered off and washed with cyclohexane to give the title compound as a yellow solid (3.75g). T.l.c. CH-EA (1:1), Rf 0.71.

Intermediate 79

20 3-Amino-1-(2-cyclopentyl-ethyl)-2,4-dioxo-5-(2-fluoro)phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

To a suspension of Zinc dust (4.70g) cooled to 0°, intermediate 78 (3.70g) in glacial acetic acid (50ml) was added. The mixture was stirred at 23° for 5h, then, diluted with water (250ml) and decanted from the zinc. Solid sodium carbonate was added until pH 9,
25 then EA(300ml) was added and the organic extracts were dried and concentrated in vacuo to give a residue which was purified by flash chromatography (eluting with CH-EA 1:1) then with DCM/methanol 9:1 to give the title compound (2.55g) as a white foam. T.l.c. DCM-methanol (90:10), Rf 0.63.

Intermediate 801-(2-cyclopentyl-ethyl)-2,4-dioxo-5-(2-fluorophenyl)-3-isocyanate-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Phosgene in toluene (1.93M solution, 25ml) was added to a solution of the intermediate
5 79 (0.734g) in dry dichloromethane (60ml); the resulting solution was stirred at 23° for
5h, then concentrated in vacuo at 50° for 3h to give the title compound as a white solid.
T.l.c. DCM-methanol (90:10), R_f 0.63.

Intermediate 812-(Bicyclo [2.2.1]-5-heptene-2-yl-methyl)-amino-diphenylamine

To a solution of 2-aminodiphenylamine (3.06g) in toluene (100ml)
5-norbornene-2-carboxaldehyde (2ml) was added and the mixture was refluxed under a
nitrogen atmosphere, in the presence of 4A molecular sieves, for 6 hrs. The solution was
decanted from the sieves and the solvent was evaporated. The residue was dissolved in
15 methanol (100ml) and sodium borohydride (5.70g) was added portionwise. The mixture
was stirred at 23° for 12 hr., diluted with ethyl acetate (100ml), washed with a 10%
potassium carbonate solution (2x100ml) and brine (100ml), then dried and concentrated
in vacuo. The crude material was purified by flash chromatography (eluting with CH-EA
95:5) to give the title compound (0.92g) as a yellow glass. T.l.c. CH-EA (95:5), R_f 0.56.

20

Intermediate 821-(Bicyclo [2.2.1]-5-heptene-2-ylmethyl)-2,4-dioxo-5-phenyl-3-phenylhydrazono-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Intermediate 81 (0.85g) and 2-phenylhydrazonomalonyl dichloride (0.87g) were each
25 taken up in dry THF (40ml) and dropped into a flask containing THF (10ml). The mixture
was refluxed, under nitrogen, for 2 hrs, then it was diluted with ethyl acetate (50ml) and
washed with a 5% sodium bicarbonate solution (50ml) and brine (50ml). The organic
layer was dried, concentrated in vacuo and purified by flash chromatography (eluting with
CH-EA 9:1) to give the title compound (1.27g) as a yellow foam. M.p. 149-151°. T.l.c.
30 (CH-EA 8:2) R_f 0.34.

Intermediate 833-Amino-1-(Bicyclo[2.2.1]-5-heptene-2-yl-methyl)-2,4-dioxo-5-phenyl-
2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Zinc dust (1.5g) was added to a solution of the intermediate 82 (0.49g) in glacial acetic acid (20ml). The mixture was stirred at 23° for 12 hrs, then it was filtered through a pad of celite. The filtrate was concentrated in vacuo; the residue was taken up in ethyl acetate (70ml) and washed with a 10% sodium hydroxide solution (2x50ml) and brine (2x50ml), then dried and concentrated in vacuo. Purification by flash chromatography (eluting with EA-MeOH 9:1) afforded the title compound (0.26g) as a light yellow foam. T.l.c.

10 (EA-MeOH 9:1), R_f 0.37.

Intermediate 843-Amino-1-(bicyclo [2.2.1]-2-heptylmethyl)-2,4-dioxo-5-phenyl-
2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

15 The intermediate 82 (0.506g), suspended in methanol (20ml), was hydrogenated at 1 atmosphere, in the presence of 5% Pd/C (0.271g) and concentrated hydrochloric acid (1.6ml), for 7h. Then, the mixture was filtered through a pad of celite and the solvents were evaporated. The residue was taken up in ethyl acetate (100ml) and washed with a 5% sodium hydroxide solution (2x100ml) and brine (100ml); the organic layer was dried, concentrated in vacuo and purified by flash chromatography (eluting with EA-MeOH 9:1) to give the title compound (0.31g) as a white foam.

20 T.l.c. (EA-MeOH 9:1) R_f 0.55.

Intermediate 852-[Bicyclo[2.2.1]-2-heptyl]amino-diphenylamine

A mixture of 2-aminodiphenylamine (5.0g), 2-norbornanone (3.0g) and molecular sieves in dry toluene (200ml) was heated to 120° for 6h. The mixture was allowed to cool to room temperature, filtered and the solution concentrated in vacuo. The residue was dissolved in ethanol (200ml), then sodium borohydride (3.0g) was added portionwise.

30 The resulting mixture was stirred at 23° for 30 min, diluted with water (150ml) and

extracted with ethyl acetate (300ml). The organic layer was washed with brine (2x200ml), dried and concentrated in vacuo to an oil which was purified by flash chromatography (eluting with CH-EA 9:1) to give the title compound as a yellow oil (3.5g). T.l.c. CH-EA (9:1), Rf 0.74.

5

Intermediate 861-[Bicyclo[2.2.1]-2-heptyl]-2,4-dioxo-5-phenylhydrazono-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

The intermediate 85 (3.77g) and the 2- phenylhydrazone malonyldichloride (3.98g) were each taken up in THF (70ml) and dropped into a flask containing THF (60ml) under a nitrogen atmosphere. After complete addition the solution was heated to 50° for 1h. The solution was concentrated in vacuo to an oil which was purified by flash chromatography (eluting with CH-EA 8:2) to give the title compound as a yellow solid (6.0g).

M.p.110-111° T.l.c. CH-EA (7:3), Rf 0.72 and 0.66.

15

Intermediate 873-Amino-1-[bicyclo[2.2.1]-2-heptyl]-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Zinc dust (3.26g) was added to a solution of the intermediate 86 (3.0g) in glacial acetic acid (30ml). The mixture was stirred at 23° for 4h, then decanted from zinc. The solution was basified until pH=9 using 10% sodium hydroxide solution and extracted with ethyl acetate (2x100ml). The combined organic extracts were washed with brine (150ml), dried and concentrated in vacuo to a residue which was triturated with diethyl ether to give the title compound as a white solid (1.34g). M.p. 172-3°. T.l.c. EA-MeOH (95:5), Rf 0.3.

25

Intermediate 882-(2-Adamantylmethyl)amino-diphenylamine

A solution of sodium acetate trihydrate (6.45g) and acetic acid (5ml) in water was added to a mixture of 2-adamantanecarboxaldehyde (2.6g) and 2-aminodiphenylamine I (2.84g) in ethanol (130ml). Then sodium borohydride (5.97g) was added portionwise. The

resulting mixture was stirred at 23° for 6h, then diluted with water (80ml) and extracted with ethyl acetate (2x150ml). The combined organic extracts were washed with brine (150ml), dried and concentrated in vacuo to a residue, which was purified by flash chromatography to give the title compound as a yellow oil (2.15g). T.l.c. CH-EA (8:2),

5 Rf 0.86.

Intermediate 89

1-(2-Adamantylmethyl)-2,4-dioxo-5-phenylhydrazono-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

10 A solution of 2-phenylhydrazonomalononyldichloride (1.78g) in THF (50ml) was added to a solution of the intermediate 88 (2.0g) in THF (50ml) under a nitrogen atmosphere. The resulting solution was heated to 50° for 1h., then concentrated in vacuo to a residue which was purified by flash chromatography (eluting with CH-EA 9:1) to give the title compound as a yellow solid (1.95g). M.p. 135- 6°(dec) T.l.c. CH-EA (8:2), Rf 0.48.

15

Intermediate 90

1-(2-Adamantylmethyl)-3-amino-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Zinc dust (1.84g) was added to a solution of the intermediate 89 (1.9g) in glacial acetic acid (20ml). The mixture was stirred at 23C for 2h, then decanted from zinc. The solution was basified until pH=9 using 10% sodium hydroxide solution and extracted with ethyl acetate (2x80ml). The combined organic extracts were washed with brine (100ml), dried and concentrated in vacuo to a residue which was purified by flash chromatography (eluting in gradient from CH-EA 1:1 to EA) to give the title compound as a yellow solid (0.95g). M.p. 209-210°. T.l.c. EA-MeOH (20:1), Rf 0.38.

Intermediate 91

5-Fluoro-N-(4-Fluorophenyl) 2-nitro aniline

A mixture of 2,4-difluoronitrobenzene (5.5ml), 4-fluoro aniline (14.2ml) and sodium carbonate (5.3g) was heated at 180° for 3h. The reaction mixture was cooled to room

temperature, then diluted with DCM , washed with water (50ml), brine (2x50ml) dried and evaporated under vacuum to give the crude compound (22.6g), which was purified by flash chromatography with CH-EA 4/1 to give the title compound as an orange solid (12.35g).M.p. 115-6° T.l.c. CH-EA (10:1), Rf 0.52.

5

Intermediate 92

4-Fluoro N¹-[4-fluorophenyl]-1,2-benzendiamine

A solution of potassium carbonate (8.292g) and sodium hydrosulfite (6.964 g) in water (200 ml) was added to a suspension of the intermediate 91 (2.502g) in 95% ethanol (350ml). The mixture was stirred at 23° for 1h, the reaction mixture was acidified to pH=3.5 with conc. hydrochloric acid and concentrated in vacuo to half volume. A 10% solution of sodium hydroxide was added until pH=10 and the solution was extracted with ethyl acetate (200ml). The combined organic extracts were washed with brine (200ml), dried and concentrated in vacuo to give the crude compound (2.93 g) which was purified by flash chromatography using CH-EA 3/2 as eluent to give the title compound as a brown oil (1.64 g). M.p.83- 84°. T.l.c. CH-EA (2:1), Rf 0.35.

Intermediate 93

N¹-(Adamantane-1-methyl)-4-Fluoro-Nⁿ-(4-fluorophenyl)-1,2- benzendiamine

To a solution of 1-adamantanecarboxaldehyde (1.223g) and intermediate 92 (1.64g) in ethanol (50ml) a buffer prepared with sodium acetate trihydrate (3.04 g) and glacial acetic acid (004ml) in water (25ml) was added and the mixture was stirred at 23° . A further amount of ethanol (15ml) was added to get a clear solution and sodium borohydride (2.8g) was added portionwise. The mixture was stirred at 23° for 20h, and then diluted with ethyl acetate (30ml). The combined organic extracts were washed with potassium carbonate (30ml) with brine (30ml), dried and concentrated in vacuo to give a red oil (3.102 g) which was purified by flash chromatography (eluting with CH-EA 15:1 to give the title compound as an orange oil (0.854g) . T.l.c. CH-EA (9:1) , Rf 0.59.

Intermediate 941-(Adamantane-1-methyl)-2,4-Dioxo-7-fluoro-5-(4-fluoro phenyl) -3-phenylhydrazone-
2,3,4,5-tetrahydro-1H-1,5-benzodiazepine.

The intermediate 93 (0.850 g) and the phenylhydrazonomalonyldichloride (0.565g) were
5 each taken up in THF (30ml) and dropped in a flask containing THF (30ml) maintained
under a nitrogen atmosphere. After complete addition the solution was heated to 70C for
3h. The solution was diluted with EA (100ml), washed with 5% sodium hydrogen
carbonate solution (100ml) and brine (100ml) , dried and concentrated in vacuo to a red
foam (1.268 g) , which was purified by flash chromatography (eluting with CH-EA 3:1)
10 to give the title compound as a yellow foam (0.562g). T.l.c. CH-EA (3:1), Rf 0.46.

Intermediate 951-(Adamantane-1-methyl)-3-Amino-2,4-dioxo-7-fluoro-5-(4-fluoro phenyl)-2,3,4,5-
tetrahydro-1H-1,5-benzodiazepine

Zinc dust (0.673g) was added to a solution of the intermediate 94 (0.557g) in glacial
15 acetic acid (20ml). The mixture was stirred at 23C for 6h, filtered and evaporated to
dryness; the residue was dissolved in water (80ml), the solution was basified with solid
sodium hydroxide until ph=9, extracted with EA (100ml). The combined organic
extracts were washed with brine (2X 30ml), dried and concentrated in vacuo to give a
20 yellow foam (0.547g) which was purified by flash chromatography (eluting with EA-
Methanol 9 /1 to give the title compound as a white solid (0.322g) M.p. 232-3°. T.l.c.
EA-methanol (9:1), Rf 0.56.

Intermediate 9625 N-1-(Adamantane-1-methyl)-3-amino-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-
benzodiazepine, (IR)-(-)-10-camphorsulphonate

A solution of (IR)-(-)-10-camphorsulphonic acid (13.2g) in acetonitrile (1035ml) was
added dropwise to a solution of intermediate 41 (33g) in acetonitile (1089ml) and the
stirred mixture was left overnight at room temperature. The precipitate was filtered and

washed with acetonitrile, (80ml) ethyl acetate (50ml) and petroleum ether (50ml) to give, after drying in vaccum, the title compound (16.17g) as a white solid . M.P. 270-2°.

Intermediate 97

5 (+)-N-1-(Adamantane-1-methyl)-3-amino-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

A suspension of intermediate 96 (6.05g) in ethyl acetate (395ml) was mixed with 5% aqueous ammonia (395ml) for 5 min, and the organic layer separated. The aqueous layer was washed with ethyl acetate (395ml) and then the ethyl acetate phase separated. The 10 combined organic extracts were dried and the solvent evaporated to give the title compound as a white foam (4.1g). T.l.c. EA-methanol (95.5), Rf 0.33 [alpha]_D = +31.

Intermediate 98

4-Fluoro-N'-(3-methylbut-1-yl)-N"-phenyl-1,2-benzendiamine

15 Bromo 3-methylbutane (0.38ml) was added to a solution of the 5-fluoro N'-phenyl-1,2-benzendiamine (0.645 g) and sodium iodide (0.476g) in dimethylformamide (25ml) under a nitrogen atmosphere. The solution was stirred at 120° for 10h, then cooled to room temperature, diluted with water (30ml) and extracted with ethyl ether (2x25ml). The combined organic extracts were washed with brine (30ml), dried and concentrated in 20 vacuo to give a red oil, which was purified by flash chromatography (eluting with CH-EA 9:1) to give the title compound as a brown oil (0.467g) . T.l.c. CH-EA (2:1) , Rf 0.78.

Intermediate 99

2,4-Dioxo-7-fluoro-1-(3-methylbut-1-yl)-5-phenyl-

25 3-phenylhydrazone-2,3,4,5-tetrahydro- 1H-1,5-benzodiazepine.

The intermediate 98 (0.454 g) and the phenylhydrazonomalonyldichloride (0.49g) were each taken up in THF (15ml) and dropped in a flask containing THF (15ml) maintained under a nitrogen atmosphere. After complete addition the solution was heated to 70° for 1h. The solution was diluted with EA (20ml), washed with 5% sodium hydrogen carbonate solution (20ml) and brine (20ml), dried and concentrated in vacuo to an oil,

which was purified by flash chromatography (eluting with CH-EA 8:2) to give the title compound as a yellow foam (0.565g). T.l.c. CH-EA (4:1), Rf 0.33.

Intermediate 100

- 5 3-Amino-2,4-dioxo-7-fluoro-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepine

Zinc dust (0.822g) was added to a solution of the intermediate 99 (0.559g) in glacial acetic acid (20ml). The mixture was stirred at 23° for 2h, then diluted with 10% solution of sodium hydroxide until pH=9 and the mixture extracted with ethyl acetate (2x30ml).

- 10 The combined organic extracts were washed with brine (30ml), dried and concentrated in vacuo to give a brown oil (0.529g) which was purified by flash chromatography (eluting with CH-Methanol 19/1 to give the title compound as a yellow foam (0.323 g). M.p. 125-6C. T.l.c. EA-methanol (19:1), Rf 0.45.

15 EXAMPLE 1

N-[2,4-Dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

- Phenyl isocyanate (0.136ml) was added to a solution of the intermediate 3 (0.4g) in dry acetonitrile (10ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, 20 filtered and the solid washed with diethyl ether to give the title compound as a white solid (0.45g). M.p. 254-50. T.l.c. CH-EA(1:1), Rf 0.65. IR :3450 (NH), 1707 and 1670 (C=O), 1601 and 1533 (C=C) cm-1; ¹H-NMR :7.459 (dd); 7.4-7.1 (m); 7.03 (m); 6.989 (dd); 6.933 (bs); 6.353 (d); 5.366 (d); 4.457 (m); 3.70 (m); 1.6-1.4 (m); 0.902 (d); 0.888 (d).

25

EXAMPLE 2

N-[1-(3,3-Dimethylbut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

- Phenyl isocyanate (0.106ml) was added to a solution of the intermediate 6 (0.3g) in 30 dry acetonitrile (5ml) under a nitrogen atmosphere. The mixture was stirred at 23° for

1h, filtered and the solid washed with diethyl ether to give the title compound as a white solid (0.27g). M.p. 271-2°. T.l.c. CH-EA (7:3), Rf 0.32. IR :3310 (NH), 1718, 1668 and 1639 (C=O), 1601 and 1556 (C=C) cm-1; ¹H-NMR :7.45 (dd); 7.4-7.10 (m); 7.06-6.97 (m); 6.414 (d); 5.362 (d); 4.476-4.373 (m); 3.757-3.656 (m); 1.503 (m); 0.924 (s).

5 EXAMPLE 3

N-[2,4-Dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-methylmercapto)phenylurea

10 3-Methylmercaptoaniline (0.065ml) was added to a solution of the intermediate 7 (0.2g) in dichloromethane (10ml) under a nitrogen atmosphere. The solution was stirred at 23° for 3h, then concentrated in vacuo and triturated with acetonitrile to give the title compound as a white solid (0.132g). M.p. 246-7°. T.l.c. CH-EA (1:1), Rf 0.58. IR: 1711, 1691, 1680 and 1670 (C=O), 1595 (C=C) cm-1; ¹H-NMR :7.46 (dd); 7.4-7.3 (m); 7.26-7.10 (m); 7.04-6.9 (m); 6.82-6.76 (bm); 6.257 (d); 5.333 (d); 4.46 (m); 3.700 (m); 2.436 (s); 1.6-1.4 (m); 0.906 (d); 0.886 (d).

15 EXAMPLE 4

N-[2,4-dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-dimethylamino)phenylurea

20 Triethylamine (0.32ml) and 3-dimethylaminoaniline dihydrochloride (0.24g) were added to a suspension of the intermediate 8 (0.22g) in dry dimethylformamide (5ml) under a nitrogen atmosphere. The resulting mixture was heated to 160° for 2h, then cooled to room temperature, diluted with water (20ml) and extracted with ethyl acetate (2x20ml).
25 The combined organic extracts were dried, concentrated in vacuo and triturated with acetonitrile to give the title compound as a white solid (0.12g). M.p. 252-3°. T.l.c. CH-EA (1:1), Rf 0.5. IR: 3312 (NH), 1707, 1676 and 1639 (C=O), 1593 and 1558 (C=C) cm-1; ¹H-NMR :7.45 (dd); 7.41-7.28 (m); 7.25-7.1 (m); 7.134 (t); 6.981 (dd); 6.818 (t); 6.634 (bs); 6.599 (dd); 6.455 (dd); 6.365 (d); 5.359 (d); 4.509-4.409 (m); 3.741
30 -3.645 (m); 2.918 (s); 1.6-1.42 (m); 0.908 (d); 0.896 (d).

EXAMPLE 5

N-[1-(3,3-Dimethylbut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-methylmercapto)phenylurea

5 3-Methylmercaptoaniline (0.19ml) was added to a solution of the intermediate 9 (0.3g) in dry dimethylformamide (5ml) under a nitrogen atmosphere. The solution was heated to 160° for 5h, then cooled to room temperature, diluted with water and extracted with ethyl acetate (2x20ml). The combined organic extracts were dried, concentrated in vacuo and triturated with acetonitrile to give the title compound as a white solid (0.08g). M.p. 10 249-50°. T.l.c. CH-EA (7:3), Rf 0.33. IR : 3308 (NH), 1707, 1676 and 1643 (C=O), 1607 (C=C) cm-1; ¹H-NMR : 7.48-7.30 (m); 7.28-7.10 (m); 7.04-6.90 (m); 6.83 (bs); 6.29 (d); 5.34 (d); 4.41 (m); 3.71 (m); 2.44 (s); 1.50 (m); 0.93 (s).

EXAMPLE 6

15 N-[1-(3,3-Dimethylbut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-dimethylamino)phenylurea
Triethylamine (0.43ml) and 3-dimethylaminoaniline dihydrochloride (0.324g) were added to a solution of the intermediate 9 (0.3g) in dry dimethylformamide (5ml) under a nitrogen atmosphere. The solution was heated to 160° for 2h, then cooled to room temperature, 20 diluted with water and extracted with ethyl acetate (2x20ml). The combined organic extracts were dried, concentrated in vacuo and triturated with acetonitrile to give the title compound as a white solid (0.16g). M.p. 255-6°. T.l.c. CH-EA (6:4), Rf 0.28. IR : 3308 (NH), 1717 (C=O), 1637 (C=C) cm-1; ¹H-NMR : 7.48-7.10 (m); 6.98 (dd); 6.81 (t); 6.66-6.56 (m); 6.46 (dd); 6.34 (d); 5.36 (d); 4.41 (m); 3.70 (m);

25

EXAMPLE 7A

N-[1-(3,3-Dimethyl-2-hydroxybut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.068ml) was added to a solution of the intermediate 12 (0.2g) in dry acetonitrile (5ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 20h,

concentrated in vacuo and the residue triturated with diethyl ether to give the title compound as a white solid (0.2g). M.p. 248-9°. T.l.c. CH-EA (1:1), Rf 0.60 and 0.58. IR: 3308 (NH), 1709 and 1670 (C=O), 1639 and 1601 (C=C) cm-1; ¹H-NMR: 7.66 (d); 7.46-7.06 (m); 7.02-6.9 (m); 6.8-6.7 (bs); 6.62 (d); 5.412 (d); and 5.402 (d); 4.492 (bd); 5 4.303 (bm); 3.936 (d); 3.95-3.85 (m); 3.613 (bt); 3.48 (bs); 2.634 (bs); 2.504 (bs); 0.918 (s).

EXAMPLE 7B

10 N-[1-(3,3-Dimethyl-2-hydroxybut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea (diastereomer I)
Phenyl isocyanate (0.0984ml) was added to a suspension of the intermediate 13a (0.29g) in dry acetonitrile (5ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 20h, concentrated in vacuo and the residue triturated with diethyl ether to give the title compound as a white solid (0.29g). M.p. 255-6° (dec). T.l.c. CH-EA (1:1), Rf 0.6. IR: 3352, 3282 and 3253 (NH and OH), 1705 and 1680 (C=O), 1630 and 1599 (C=C) cm-1; 15 ¹H-NMR: 7.905 (dd); 7.38-7.24 (m); 7.24-7.1 (m); 7.05 (bs); 6.98-6.85 (m); 6.80 (bs); 5.395 (d); 4.513 (bd); 3.936 (bs); 3.598 (bt); 2.521 (bs); 0.924 (s).

EXAMPLE 7C

20 N-[1-(3,3-Dimethyl-2-hydroxybut-1-yl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea (diastereomer II)
Phenyl isocyanate (0.12ml) was added to a suspension of the intermediate 13b (0.33g) in dry acetonitrile (5ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 3h, filtered and the solid washed with diethyl ether to give the title compound as a white solid (0.27g). M.p. 204-5°. T.l.c. CH-EA (1:1), Rf 0.58. IR: 3308 (NH and OH), 1718 and 1670 (C=O), 1601 (C=C) cm-1; ¹H-NMR: 7.86 (d); 7.4-7.12 (m); 7.02-6.94 (m); 25 6.577 (d); 5.414 (d); 4.312 (t); 3.931 (d); 3.454 (bs); 2.560 (bs); 0.919 (s).

EXAMPLE 8

N-[1-(1,3-Dimethylbut-1-yl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.1ml) was added to a solution of the intermediate 16 (0.22g) in dry acetonitrile (10ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, 5 concentrated in vacuo to give an oil which was purified by flash chromatography (eluting with CH-EA 80:20 to give a crude sample which was triturated with 1/1 mixture of petroleum/ethyl ether (30 ml) to give the title compound (0.12 g). T.l.c. CH-EA(1:1), Rf 0.53. IR :3370 (NH), 1701 and 1670 (C=O), 1651 and 1601 (C=C) cm-1; ¹H-NMR :7.44-7.35 (m); 7.34-7.24 (m); 7.24-7.15 (m); 6.982 (m); 6.538 (d); 10 6.529 (d); 5.328 (d); 5.321(d); 4.576 (m); 4.438(q); 2.11(m); 1.74-1.64(m); 1.64-1.44(m); 1.542 (d); 1.435(d); 0.886 (d); 0.882 (d); 0.873 (d); 0.827(d).

EXAMPLE 9

N-[7-Chloro-2,4-dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.1ml) was added to a solution of the intermediate 20 (0.2g) in dry acetonitrile (9ml) under a nitrogen atmosphere. The mixture was stirred at 0° for 2h, filtered and the solid triturated with petroleum ether/ethyl ether (2/2ml) at 0°C, filtered off, washed with 1/1 mixture petroleum ether/ethyl ether (10 ml) to give the title compound as a white solid (0.17g). T.l.c. CH-EA(1:1), Rf 0.59. IR :3312 (NH), 1713 and 1684 (C=O), 1639 and 1605 (C=C) cm-1; ¹H-NMR :7.45-7.00 (m); 7.10(m); 6.989 (dd); 6.97 (d); 6.42 (d); 5.31 (d); 4.51(m); 3.59(m); 1.58-1.46 (m); 1.46-1.38(m); 0.87 (d); 0.85(d).

25 EXAMPLE 10

N-[8-Chloro-2,4-dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.1ml) was added to a solution of the intermediate 25 (0.2g) in dry acetonitrile (4ml) under a nitrogen atmosphere. The mixture was stirred at 0° for 30min, 30 then petroleum ether was added and stirring was continued for 1h. the solid was filtered

off, washed with 3/1 mixture petroleum ether/ethyl ether (15 ml) to give the title compound as a white solid (0.22g). T.l.c. CH-EA(1:1), Rf 0.63. IR :3310 (NH), 1717, 1668 and 1641 (C=O), cm-1; ¹H-NMR :7.44-7.35 (m); 7.32(t); 7.25-7.16 (m); 7.14 (m); 7.03 (m); 6.92 (d); 6.41(d); 5.31 (d); 4.52(m); 3.62(m); 1.60- 1.40 (m); 0.89(d); 5 0.87(d).

EXAMPLE 11

N-[7,8-dichloro-2,4-Dioxo-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

10 Phenyl isocyanate (0.09ml) was added to a solution of the intermediate 30 (0.19g) in dry acetonitrile (2.5ml) under a nitrogen atmosphere. The mixture was stirred at 0° for 20min, then evaporated to dryness and the resulting solid solid triturated with 1/1 mixture petroleum ether/ethyl ether (10ml) at 0°C for 1h, filtered off, washed with 1/1 mixture petroleum ether/ethyl ether (15 ml) to give the title compound as a white solid 15 (0.15g). T.l.c. CH-EA(1:1), Rf 0.6. IR :3375 (NH), 1711, 1684 and 1655 (C=O), 1599, 1547 9 C=C) cm-1; ¹H-NMR :7.51 (s); 7.46-7.32(m); 7.28- 7.14 (m); 7.05 (s); 7.06-7.00 (m); 6.40 (d); 5.31 (d); 4.50m); 3.56(m); 1.60-1.40 (m); 0.89(d); 0.86(d).

EXAMPLE 12

N-[2,4-Dioxo-8-Fluoro--1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

20 Phenyl isocyanate (0.1ml) was added to a solution of the intermediate 35 (0.2g) in dry acetonitrile (2.5ml) under a nitrogen atmosphere. The mixture was stirred at 0° for 30min, then diethyl ether (5 ml) was added and stirring continued for 1h. The resulting 25 solid was filtered off, washed with 1/1 mixture petroleum ether/ethyl ether (10ml) to give the title compound as a white solid (0.25g). T.l.c. CH-EA(1:1), Rf 0.53. IR :3312(NH), 1718, 1670(C=O), 1639, 1605 (C=C) cm-1; ¹H-NMR :7.44-7.36 (m); 7.32(t); 7.30-7.10(m); 7.06-6.9 (m); 6.35(d); 5.33 (d); 4.52(m); 3.62(m); 1.60-1.40 (m); 0.90(d); 0.87(d).

EXAMPLE 13N-[2,4-Dioxo-5-phenyl-1-(2-phenylethyl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.1ml) was added to a solution of the intermediate 38 (0.3g) in dry acetonitrile (15ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, EE (30ml) was added and the formed precipitate was stirred for 45 min at 0°. The precipitate was filtered, washed with diethyl ether (25ml) to give the title compound as a white solid (0.27g). T.l.c. CH-EA (1:1), $R_f=0.45$. IR: 3310 (NH), 1707, 1678 (C=O); 1643, 1603, 1556 (C=C) cm^{-1} ; $^1\text{H-NMR}$: 7.428 (dd), 7.36-7.27 (m), 7.27-7.12 (m), 10 7.07-6.94 (m), 6.484 (d), 5.361 (d), 4.78-4.66 (m), 3.98-3.86 (m), 2.927 (m).

EXAMPLE 14N-[1-(1-Adamantyl)methyl-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

15 Phenyl isocyanate (0.039ml) was added to a solution of the intermediate 41 (0.13g) in dry acetonitrile (7ml) under a nitrogen atmosphere. The mixture was stirred for 1.5 hrs and the formed precipitate was filtered washing with acetonitrile (3ml) to give the title compound as a white solid (0.085g). T.l.c. CH-EA (1:1), $R_f=0.23$. IR: 3294 (NH), 1717, 1705, 1680 (C=O); 1643 (C=C) cm^{-1} ; $^1\text{H-NMR}$: 7.5-6.96 (m), 7.08 (bs), 6.50 (d), 5.31 (d), 4.49 (d), 3.37 (d), 1.84 (m), 1.6-1.3 (m).

EXAMPLE 15N-[1-(2,2-Dimethylethoxycarbonylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

25 Phenyl isocyanate (0.091ml) was added to a solution of the intermediate 44 (0.244g) in dry acetonitrile (16ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 2h, dichloromethane (30ml) was added and the organic layer was washed with brine (2x10ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with DCM-methanol 98:2), to give the title compound as a

white solid (0.232g). T.l.c. DCM- methanol (95:5), $R_f=0.8$. IR: 3431, 3395 (NH), 1745, 1684 (C=O) cm^{-1} ; $^1\text{H-NMR}$: 7.4-6.95 (m), 7.1 (bs), 6.5 (d), 5.45 (d), 4.61 (dd), 1.40 (s).

EXAMPLE 16

5 N-[1-(3,3-Dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.067ml) was added to a solution of the intermediate 47 (0.190g) in dry acetonitrile (10ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h and the formed precipitate was filtered washing with acetonitrile (3ml), to give the title compound as a white solid (0.198g). T.l.c. CH-EA (1:1), $R_f=0.57$. IR: 3431, 3350 (NH), 1745, 1668 (C=O) 1599 (C=C) cm^{-1} ; $^1\text{H-NMR}$: 7.48-7.26 (m), 7.26-7.14 (m), 7.04-6.96 (m), 6.523 (d), 5.352 (d), 4.511-4.409 (m), 1.467 (t), 0.915 (s).

EXAMPLE 17

15 N-[2,4-Dioxo-1-(2-hydroxy-3-methylbutyl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

To a solution of intermediate 63 (0.12g) in methanol (20ml) and water (3ml), sodium boro hydride (1.5g) was added portionwise at 0° , maintaining the pH at 7-7.5 by adding a 1M solution of hydrochloric acid. During the reaction, further methanol was added. The reaction mixture was stirred for 1 hr, then concentrated, diluted with ethyl acetate (100ml) and washed with brine (3x70ml), dried and concentrated "in vacuo". The crude product was dissolved in diethyl ether (5ml) and precipitated with petroleum ether (10ml), to give the title compound as a white solid (0.07g). T.l.c. CH-EA (1:1), $R_f=0.36$. IR: 3337 (NH, OH), 1701, 1647 (C=O); 1597, 1553 (C=C) cm^{-1} ; $^1\text{H-NMR}$: 7.6-6.65 (m), 5.37 (d), 5.35 (d), 3.92 (bm), 3.48 (bm), 4.50 (dd), 3.80 (dd), 4.34 (dd), 3.57 (dd), 2.50 (bm), 1.58 (m), 0.93- 0.87 (m).

EXAMPLE 18

30 N-[1-(3,3-Dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-trifluoromethoxyphenyl)urea

A solution of 3-trifluoromethoxyphenylamine (0.047g) and intermediate 48 (0.100g) in dichloromethane (5ml) were stirred for 20h at 23° under a nitrogen atmosphere, then concentrated "in vacuo". The crude product was triturated with acetonitrile (2ml) to obtain the title compound as a white solid (0.067g). T.l.c. CH-EA (60:40), $R_f=0.57$. IR: 5 3317 (NH), 1717, 1650 (C=O); 1609, 1558 (C=C) cm^{-1} ; $^1\text{H-NMR}$: 7.53 (bs), 7.46 (dd), 7.45-7.30 (m), 7.30-7.18 (m), 7.10 (t), 7.00 (dd), 6.88 (m), 6.77 (m), 6.66 (d), 5.35 (d), 4.44 (m), 3.70 (m), 1.54-1.42 (m), 0.91 (s).

EXAMPLE 19

10 N-[1-(3,3-Dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-cyanophenyl)urea

A solution of 3-cyanophenylamine (0.118g) and intermediate 48 (0.339g) in dichloromethane (10ml) were stirred for 5h at 23° under a nitrogen atmosphere, then concentrated "in vacuo". The crude product was triturated with acetonitrile (8ml), filtered and washed with acetonitrile (3ml) to obtain the title compound as a white solid (0.216g). T.l.c. CH-EA (1:1), $R_f=0.55$. IR: 3319 (NH), 2230 (C=N), 1711, 1647 (C=O); cm^{-1} ; $^1\text{H-NMR}$: 7.91 (bs); 7.52-7.30 (m), 7.30-7.12 (m), 7.01 (dd), 6.88 (d), 5.34 (d), 4.52-4.38 (m), 3.80-3.68 (m), 1.51 (m), 0.91 (s).

20 EXAMPLE 20

N-[1-(3,3-Dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-methylthiophenyl)urea

To a solution of the intermediate 49 (0.20g) in dry DMF (5ml), 3- methylthiophenylamine (0.218ml) was added and the reaction mixture was stirred for 4h at 120°, under a nitrogen atmosphere. Ethyl acetate (50ml) added and the solution washed with water (2x25ml), and brine (25ml), dried and concentrated "in vacuo". The crude product was triturated with acetonitrile (4ml) to obtain the title compound as a white solid (0.115g). T.l.c. CH-EA (1:1), $R_f=0.62$. IR: 3300 (NH), 1705, 1674, 1641 (C=O); 1607 (C=C) cm^{-1} ; $^1\text{H-NMR}$: 7.48-7.10 (m), 7.02-6.90 (m), 6.82 (s), 6.30 (d), 5.30 (d), 4.46 (m), 3.70 (m), 30 2.44 (s), 1.48 (t), 0.93 (s).

EXAMPLE 21

N-[1-(3,3-Dimethylbutyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-N,N-dimethylaminophenyl)urea

5 To a solution of the intermediate 49 (0.20g) in dry DMF (8ml), 3-N,N-dimethylaminophenylamine hydrochloride (0.177g) and triethylamine (0.118ml) were added and the reaction mixture was stirred for 4h at 120°, under a nitrogen atmosphere. Ethyl acetate (50ml) added and the solution washed with water (2x25ml) and brine (25ml), dried and concentrated "in vacuo". The crude product was purified by flash chromatography (eluting with CH-EA 60:40), then triturated with a mixture of ethyl acetate and petroleum ether to obtain the title compound as a white solid (0.076g). T.l.c. CH-EA (1:1), R_f =0.31. IR: 3500 (NH), 1794, 1707, 1666 (C=O); 1607 (C=C) cm⁻¹; ¹H-NMR: 7.46-7.10 (m), 6.99 (dd), 6.82 (t), 6.60 (m), 6.46 (m), 6.53 (bs), 6.31 (d), 5.31 (d), 4.47 (m), 3.69 (m), 2.94 (s), 2.93 (s), 1.47 (m), 0.94 (s).

15

EXAMPLE 22

N-[1-[2-(1-Adamantyl)ethyl]-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-N,N-dimethylaminophenyl)urea

To a solution of the intermediate 50 (0.12g) in dry DMF (2ml), 3-N,N-dimethylaminophenylamine dihydrochloride (0.084g) and triethylamine (0.1ml) were added and the reaction mixture was stirred for 9h at 120°, under a nitrogen atmosphere. Ethyl acetate (50ml) added and the solution washed with a saturated solution of ammonium chloride (50ml) and brine (3x50ml), dried and concentrated "in vacuo". The crude product was triturated with acetonitrile (10ml) to obtain the title compound as a white solid (0.030g). T.l.c. CH-EA (1:1), R_f =0.37. IR: 3373 (NH), 1707, 1682, 1660 (C=O); 1595 1580 (C=C) cm⁻¹; ¹H-NMR: 7.45-6.35 (m), 7.34-7.26 (m), 7.22-7.15 (m), 7.116 (t), 6.978 (dd), 6.740 (bs), 6.563 (dd), 6.44 (dd), 6.418 (d), 5.314 (d), 4.523-4.420 (m), 3.721-3.621 (m), 2.911 (s), 1.936 (bs), 1.672 (bq), 1.500 (d), 1.332 (t).

30

EXAMPLE 23

N-[1-(2,3-Dimethyl)butyl-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.03ml) was added to a solution of the intermediate 56 (0.087g) in dry acetonitrile (3ml), at 0°, under a nitrogen atmosphere. The mixture was allowed to stand at 23° and stirred for 1h, then petroleum ether was added and the formed precipitate was stirred for 4h, filtered and washed with petroleum ether. The precipitate was triturated with a mixture petroleum- ether/diethyl ether (1:1; 10ml) for 1 hr and filtered to give the title compound as a white solid (0.08g). T.l.c. CH-EA (1:1), R_f=0.49. IR: 3300 (NH), 1707, 1641 (C=O); 1558, 1541 (C=C) cm⁻¹; ¹H-NMR: 7.46-7.10 (m), 6.9 (m), 6.4 (m), 5.32 (d), 5.29 (d), 4.61 (dd), 4.48 (dd), 3.60 (dd), 3.42 (dd), 1.8 (m), 1.4 (m), 0.86 (d), 0.80 (d), 0.77 (d), 0.75 (d), 0.73 (d), 0.70 (d).

EXAMPLE 24

N-[1-Butyl-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.04ml) was added to a solution of the intermediate 59 (0.09g) in dry acetonitrile (10ml), at 23°, under a nitrogen atmosphere. The mixture was stirred for 3h, dichloromethane (30ml) added and the solution washed with water (50ml), purified by filtration on a pad of silica (eluting with DCM), to give the title compound as a white solid (0.1g). T.l.c. DCM-methanol (95:5), R_f= 0.65. IR: 3431 (NH), 1707, 1670 (C=O); 1599 (C=C) cm⁻¹; ¹H-NMR: 7.4- 7.00 (m), 6.66 (bs), 6.22 (d), 5.3 (d), 4.55 (m), 3.7 (m), 1.53 (m), 1.3 (m), 0.88 (t).

EXAMPLE 25

N-[2,4-Dioxo-5-phenyl-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.08ml) was added to a solution of intermediate 66 (0.206g) in dry acetonitrile (12ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, then dichloromethane was added until complete dissolution of the precipitate. The organic layer was separated, washed with brine (3x20ml) dried and concentrated in vacuo to give

the crude compound (0.3g) which was purified by flash chromatography (eluting with DCM-methanol 98:2) to give the title compound as a white solid (0.06g). T.l.c. DCM-methanol (95:5), Rf 0.87. IR : 3440-3350 (NH), 1701 and 1680 (C=O), 1616 and 1599 (C=C) cm⁻¹; ¹H-NMR : 7.44-7.16 (m); 7.00 (m); 6.4(m); 5.33(d); 4.53(m); 3.68 (m); 5 1.6-1.4 (m); 0.89 (d); 0.86(d).

EXAMPLE 26

(+)-N-[2,4-Dioxo-5-phenyl-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

10 Phenyl isocyanate (0.15ml) was added to a solution of intermediate 70 (0.42g) in dry acetonitrile (20ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, then the precipitate was filtered off and washed with acetonitrile (10 ml) and dried to give the title compound as a white solid (0.52g). [alpha]_D=+116. T.l.c. DCM-methanol (95:5), Rf 0.87. IR : 3308 (NH); 1703-1674 (C=O), 1645 and 1601(C=C) cm⁻¹; ¹H-NMR : 7.5-7.1 (m); 6.98 (m); 6.58(d); 5.34(d); 4.53(m); 3.68 (m); 1.58-1.4 (m); 0.87 (d); 0.84(d).

EXAMPLE 27

(+)-N-[2,4-Dioxo-5-(2-fluorophenyl)-1-(3-methylbut-1-yl)-2,3,4,5-tetrahydro-1H-

20 1,5-benzodiazepin-3-yl]-N'-[3-(N,N-dimethylamino)phenyl]urea

3-(N,N-Dimethylamino)phenyl isocyanate (0.257g) was added to a solution of the intermediate 72 (0.47g) in dry acetonitrile (10ml) under a nitrogen atmosphere. The reaction mixture was stirred at 23° for 1h and the formed precipitate was filtered to give the title compound as a white solid (0.58g) in enantiomeric ratio (+)/(-)=93/7. A sample was purified by HPLC to give the pure title compound. M.p. 252-3. T.l.c. CH-EA (1:1), Rf 0.50. [alpha]_D=+109.6. IR (nujol): 3420 (NH), 1717, 1701, 1690 and 1649 (C=O), 1616 and 1560 (C=C) cm⁻¹; ¹H-NMR : 7.45 (dd); 7.42-7.28 (m); 7.25-7.1 (m); 6.98 (dd); 6.82 (t); 6.60 (m); 6.45 (dd); 6.356 (d); 5.36 (d); 4.52-4.38 (m); 3.80-3.60 (m); 2.92 (s); 1.66-1.4 (m); 0.90 (d); 0.89 (d).

EXAMPLE 28

N-[1-(Adamant-2-yl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N-[3-(N,N-dimethylaminophenyl)]urea

Triethylamine (0.065ml) and 3-dimethylaminoaniline dihydrochloride (0.049g) were
5 added to a solution of the intermediate 76 (0.1g) in dichloromethane (5ml) under a
nitrogen atmosphere. The solution was stirred at 23 for 3h, then concentrated in vacuo
and purified by flash chromatography (eluting with CH-EA 1:1) to give the title
compound (0.052g) as a white solid. T.l.c. DCM-methanol (95:0.5), Rf 0.72. IR : 3300
(NH), 1713 and 1676 (C=O), 1637 and 1610 (C=C) cm- 1; 1H-NMR : 7.4-7.1 (m); 6.99
10 (m); 6.80 (t); 6.62 (m); 6.56 (dd); 6.45 (dd); 6.31 (d); 5.31 (d); 4.52 (m); 2.91 (m); 2.32
(m); 2.0-1.1 (m).

EXAMPLE 29

N-(2-cyclopentyl-ethyl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-
15 benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.044ml) was added to a solution of the intermediate 79 (0.154g) in
acetonitrile (5ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h; the
obtained solid was filtered and washed with acetonitrile (2 ml) to give the the title
compound (0.163g) as a white solid. Mp 255-257° T.l.c.CH-EA (1:1), Rf 0.58 . IR :
20 3400 (NH), 1718 and 1650 (C=O), 1600 (C=C) cm-1; 1H-NMR : 7.46 (dd); 7.4-7.1(m);
7.0(t); 6.98 (d); 6.52 (d); 5.38 (d); 4.44(m); 3.66 (m); 1.84-1.40 (m); 1.20-1.00 (m).

EXAMPLE 30

N-[1-(2-cyclopentyl-ethyl)-2,4-dioxo-5-(2-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5-
25 benzodiazepin-3-yl]-N'-[4-(dimethylamino)phenyl]urea

Triethylamine (0.184ml) and 4-(dimethylamino)aniline (0.138g) were added to a solution
of the intermediate 80 (0.270g) in dry dichloromethane (50ml) under a nitrogen
atmosphere. The solution was stirred at 23° for 4h, then diluted with dichloromethane
(20ml) and washed with water (20ml), 5% HCl solution (20ml), water (20ml)and brine
30 (15ml). The organic layer was dried, concentrated in vacuo, and the residue was purified

by flash chromatography (eluting with DCM-methanol 95:5) to give the title compound (0.077g) as a white solid. T.l.c. DCM-methanol (9:1), R_f 0.81. IR : 3304 (NH), 1718-1641 (C=O), 1605-1549 (C=C) cm⁻¹; 1H-NMR : 7.46(dd); 7.40-7.10 (m); 6.98 (dd); 6.68 (d); 6.28 (bs); 6.07 (d); 5.32 (d); 4.41 (m); 3.66 (m); 2.91 (s); 1.84-1.00 (m).

5

EXAMPLE 31

N-[1-(Bicyclo [2.2.1]-5-heptene-2-ylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

10 Phenylisocyanate (0.026ml) was added to a solution of the intermediate 83 (0.074g) in dry acetonitrile (5ml) and the mixture was stirred at 23°, under nitrogen, for 1 h. Dichloromethane (50ml) was added until complete dissolution of the precipitate, then the solution was washed with brine (20ml), dried and concentrated in vacuo. The residue was triturated with diethyl ether to give the title compound (0.0521g) as an off-white solid.

15 M.p. 184-6 C. T.l.c. (CH-EA 7:3) R_f 0.32. IR:3308 (NH), 1715-1670 (C=O), 1639-1599 (C=C); 1H-NMR: 7.5-7.0 (m); 6.84 (bs); 6.80 (bs); 6.33(d); 6.31 (d); 6.18-6.10 (m); 6.12-5.96 (m); 5.90-5.84 (m); 5.64-5.60 (m); 5.32 (d); 5.29 (d); 4.64 (m); 4.4-4.2 (m); 3.8 (m); 3.45-3.30 (m); 2.80 (bs); 2.74 (bs); 2.6-0.60 (m).

20

EXAMPLE 32

N-[1-(Bicyclo [2.2.1]-5-heptene-2-ylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-nitrophenyl)-urea

To a solution of the intermediate 83 (0.072g) in dry acetonitrile (10ml) 3-nitrophenylisocyanate (0.048g) was added and the mixture was stirred at 23°, under nitrogen, for 1 h. The resulting precipitate was filtered, washed with diethyl ether, and dried to give the title compound (0.0712g). M.p. 195-7 C.

T.l.c. (CH-EA 7:3) R_f 0.24. IR : 3300 (NH); 1713 (C=O); 1651 (C=O); 1556 (C=C)cm⁻¹. 1H-NMR : 8.26-6.92 (m); 6.13- 5.58 (m); 5.34-5.25 (m); 4.70-3.83 (m); 2.80-0.45 (m).

30

EXAMPLE 33N-[1-(Bicyclo [2.2.1]-2-heptylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3yl]-N'-phenylurea

Phenylisocyanate (0.03ml) was added to a solution of the intermediate 84 (0.088g) in dry acetonitrile (10ml) and the mixture was stirred at 23°, under nitrogen, for 1 h. The resulting precipitate was filtered, washed with diethyl ether and dried to give the title compound (0.0858g) as a white solid. M.p. 255-6°. T.l.c. (CH-EA 7:3) R_f 0.29. IR: 3400-3200 (NH), 1711 and 1705 (C=C); 1H-NMR: 7.50-7.10 (m), 7.02 (m), 6.38 (m), 6.91 (bs), 6.42-6.34 (m), 5.35-5.27(m), 4.71-4.61 (dd), 4.48 (dd), 4.38 (dd), 3.65 (dd), 3.59 (dd), 3.37 (dd), 3.55 (dd), 2.3-0.50 (m).

EXAMPLE 34N-[1-(Bicyclo [2.2.1]-2-heptylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3yl]-N'-(3-methoxyphenylurea)

To a solution of the intermediate 84 (0.0883g) in dry acetonitrile (10ml) and the mixture was stirred at 23° under nitrogen, for 1h. The resulting precipitate was filtered, washed with diethyl ether and dried to give the title compound (0.0858g) as a white solid. M.p. 255-6°. T.l.c. (CH-EA 7:3) R_f 0.29. IR: 3400-3200(NH), 1711 and 1705 (C=C); ¹H-NMR: 7.5-6.9(m); 6.75-6.69(m); 6.60-6.55(m); 6.44(m); 5.31(m); 4.71-4.62(m); 4.49(dd) 4.38(dd); 3.75(s); 3.65(dd); 3.59(dd); 3.56(dd); 3.38(dd); 2.25-0.6(m).

Example 35N-[1-[Bicyclo[2.2.1]-2-heptyl]-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.056ml) was added to a solution of the intermediate 87 (0.15g) in dry acetonitrile (5ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, then filtered. The solid obtained was washed with diethyl ether and dried in vacuo to give the title compound as a white solid (0.12g). M.p. 267-8°. T.l.c. CH-EA(1:1), R_f 0.62. IR: 3300 (NH), 1705, 1678 and 1645 (C=O), 1599 and 1556 (C=C) cm⁻¹; ¹H-NMR: 7.46-7.12 (m); 7.026-6.94 (m); 6.423 (d); 6.436 (d); 5.328 (d); 5.321 (d); 4.5-4.4 (m);

3.459 (s); 2.637 (s); 2.396 (m); 2.180 (m); 1.958 (m); 1.6 (m); 1.54-1.38 (m); 1.38-1.1 (m); 0.99 (m); 0.864 (m).

Example 36

5 N-[1-(2-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyanate (0.063ml) was added to a solution of the intermediate 90 (0.2g) in dry acetonitrile (5ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, then filtered. The solid obtained was washed with diethyl ether and dried in vacuo to give 10 the title compound as a white solid (0.22g). M.p. 192-3°. T.l.c. CH-EA (1:1), R_f 0.73. IR :3306 (NH), 1717 and 1701 (C=O), 1643 and 1620 (C=C) cm⁻¹; ¹H-NMR :7.5-7.14 (m); 7.00 (m); 7.049 (m); 6.47 (d); 5.33 (d); 5.05 (m); 3.59 (m); 2.02 (m); 1.84-1.36 (m).

15 EXAMPLE 37

5 N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-methoxyphenyl)urea

3-Methoxyphenyl isocyanate (0.066ml) was added to a solution of the intermediate 41 (0.2g) in dry acetonitrile (10ml) under a nitrogen atmosphere. The mixture was stirred at 20 23° for 16h, then diluted with dichloromethane (15ml) and washed with brine (15ml). The organic solution was dried, concentrated in vacuo and the residue was purified by flash chromatography (eluting with CH-EA 2:1); the solid obtained was further purified by trituration with diethyl ether to give the title compound as a white solid (0.2g). M.p. 267-8°. T.l.c. CH-EA (2:1), R_f 0.2. IR :3302 (NH), 1713, 1674 and 1641 (C=O), 1612 and 1558 (C=C) cm⁻¹; ¹H-NMR :7.492 (dd); 7.45-7.35 (m); 7.35-7.25 (m); 7.162 (m); 7.120(t); 7.041 (t); 6.992 (dd); 6.904 (s); 6.738(m); 6.578 (m); 6.413 (m); 5.292 (d); 4.496 (d); 3.744 (s); 3.382 (d); 1.857 (s); 1.66-1.32 (m).

EXAMPLE 38

N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-methylphenyl)urea

3-Methylphenyl isocyanate (0.064ml) was added to a solution of the intermediate 41 (0.2g) in dry acetonitrile (10ml) under a nitrogen atmosphere. The mixture was stirred at 5 23° for 1h, then diluted with dichloromethane (15ml) and washed with brine (15ml). The organic solution was dried, concentrated in vacuo and the residue was triturated with diethyl ether to give the title compound as a white solid (0.2g). M.p. 244-6 $^{\circ}$. T.l.c. CH-EA (2:1), Rf 0.32. IR :3300 (NH), 1715 and 1672 (C=O), 1645 and 1616 (C=C) cm-1; 1H-NMR :7.493 (dd); 7.45-7.35 (m); 7.35-7.25 (m); 7.21-7.15 (m); 7.135 (t); 10 7.034 (m); 6.992 (dd); 6.854 (m); 6.751 (s); 6.318 (d); 5.293 (d); 4.498 (d); 3.384 (d); 2.287 (s); 1.865 (s); 1.68-1.3 (m).

EXAMPLE 39

N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-nitrophenyl)urea

A solution of 3-nitrophenyl isocyanate (0.082g) in dry acetonitrile (8ml) was added to a solution of the intermediate 41 (0.2g) in dry acetonitrile (10ml) under a nitrogen atmosphere. The mixture was stirred at 20 23° for 2h, then diluted with dichloromethane (15ml) and washed with brine (15ml). The organic solution was dried, concentrated in vacuo and the residue was triturated with diethyl ether to give the title compound as a white solid (0.229g). M.p. 213-5 $^{\circ}$. T.l.c. CH-EA (2:1), Rf 0.33. IR :3296 (NH), 1713 and 1645 (C=O), 1597 (C=C) cm-1; 1H-NMR :8.25 (s); 8.15 (t); 7.64 (m); 7.52 (dd); 7.45 (m); 7.36-7.29 (m); 7.24-7.17 (m); 7.13 (t); 7.06 (d); 7.02 (dd); 5.27 (d); 4.51 (d); 3.40 (d); 1.86 (s); 1.66-1.34 (m).

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EXAMPLE 40

N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-bromophenyl)urea

3-Bromophenyl isocyanate (0.063ml) was added to a solution of the intermediate 41 (0.2g) in dry acetonitrile (10ml) under a nitrogen atmosphere. The mixture was stirred at 30

23° for 1h, then filtered. The solid obtained was washed with diethyl ether to give the title compound as a white solid (0.25g). M.p. 254-6°. T.l.c. CH-EA (2:1), Rf 0.53. IR :3290 (NH), 1717 and 1672 (C=O) cm-1; 1H-NMR :7.56-7.15 (m); 7.03-6.88 (m); 6.99 (dd); 6.93 (dd); 6.73 (d); 5.29 (d); 4.49-3.38 (m); 1.83 (m); 1.64-1.30 (m).

5

EXAMPLE 41

N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3- yl]-N'-(3-ethoxycarbonylphenyl)urea

3-Nitrophenyl isocyanate (0.1g) was added to a solution of the intermediate 41 (0.415g) 10 in dry acetonitrile (13ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, then diluted with dichloromethane (20ml) and washed with brine (20ml). The organic solution was dried, concentrated in vacuo and the residue was triturated with diethyl ether to give the title compound as a white solid (0.407g). M.p. 246-8°. T.l.c. CH-EA (2:1), Rf 0.37. IR :1709, 1690 and 1670 (C=O) cm-1; 1H-NMR : 7.93 (t); 7.64-7.50 (m); 7.44-7.39 (m); 7.38 (s); 7.35-7.27 (m); 7.24- 7.14 (m); 6.89 (dd); 6.58 (d); 5.31 (d); 4.50 (d); 4.34 (m); 3.38 (d); 1.85 (m); 1.61-1.51 (m); 1.45-1.37 (m); 1.35 (t).

EXAMPLE 42

N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3- yl]-N'-[3-(N,N-dimethylamino)phenyl]urea

A solution of 3-(N,N-dimethylamino)phenyl isocyanate (0.122g) in dry acetonitrile (7ml) 20 was added to a solution of the intermediate 41 (0.2g) in dry acetonitrile (7ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 30min, then diluted with dichloromethane (20ml) and washed with brine (20ml). The organic solution was dried, concentrated in vacuo and the residue was triturated with diethyl ether to give the title compound as a white solid (0.221g). M.p. 263-5°. T.l.c. CH-EA (1:1), Rf 0.52. IR :3300 (NH), 1717 and 1674 (C=O) cm-1; 1H-NMR : 7.48 (dd); 7.45- 7.24 (m); 7.19-7.10 (m); 6.98 (dd); 6.93 (dd); 6.61 (s); 6.58-6.45 (m); 6.38 (d); 5.29 (d); 4.49-3.37 (m); 2.92 (s); 1.87 (m); 1.63- 1.53 (m); 1.44-1.34 (m).

EXAMPLE 43N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-carboxyphenyl)urea

An aqueous 0.1M solution of lithium hydroxide (6.6ml) was added to a solution of Example 41 (0.2g) in THF (15ml) previously cooled to 0°. The solution was stirred at 23° for 16h, then heated to 60° for 1h and to 80° for 13h. The solution was cooled to 23°, neutralized with acetic acid, concentrated in vacuo and the residue purified by flash chromatography (eluting in gradient from CH-EA 3:1 to DCM and finally to DCM-MeOH 10:1) to give the title compound as a white solid (0.183g), still containing traces of inorganic salts. A sample was further purified by dissolution in DCM and washing with 10% hydrochloric acid; the organic layer was dried, concentrated in vacuo and the residue triturated with diethyl ether to give the pure title compound. M.p. 260-70°(dec). T.l.c. EA, Rf 0.64. IR: 3354 (NH and OH), m 1701 and 1684 (C=O) cm⁻¹; ¹H-NMR: 9.21(s); 7.9-7.8(m); 7.6-7.16(m); 7.0-6.9(m); 4.99(d); 4.30(d); 3.60(d); 1.83(s); 1.65-1.2(m).

EXAMPLE 44N-[1-(Adamantanmethyl)-2,4-dioxo-7-fluoro-5(4-fluorophenyl)-2,3,4,5-tetrahydro-1H-1,5benzodiazepin-3-yl]-N'-(3-dimethylamino)phenylurea

A solution of 3-Dimethylaminophenyl isocyante (0.043g) in dry acetonitrile (3ml) was added to a solution of the intermediate 95 (0.079g) in dry acetonitirle (5ml) under a nitrogen to a solution of the mixture was stirred at 23C for 1h diluted with DCM, washed with brine (30ml), evaporated to give the crude compound (0.145g) which was triturated with ethyl ether to give the title compound as a white solid (0.046g) M.p. >270. T.l.c. CH-EA (1:1), Rf 0.61. IR: 3439, 3333(NH); 1715 (C=O), 1610 and 1590 (C=C) cm⁻¹; ¹H-NMR: 7.46(dd); 7.38-7.3(m); 7.20-7.10(m); 7.06-7.00(m); 6.78(t); 6.69-6.58(m); 6.49(dd); 6.27(d); 5.26(d); 4.49(d); 3.28(d); 2.93(s); 1.8°(bs); 1.67-1.30(m).

EXAMPLE 45(+)-N-[1-(Adamantanmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5benzodiazepin-3-yl]-N'-phenylurea

Phenyl isocyante (0.033ml) was added to a solution of intermediate 97 (0.096g) in dry acetonitrile (9ml). The mixture was stirred at 23° under a nitrogen atmosphere for 1h, then it was diluted with dichloromethane (40ml) and washed with brine (2x20ml). The organic layer was dried and concentrated in vacuo. Crystallisation of the crude material from ethyl acetate afforded the title compound (0.075g) as white needles. M.P. 264-5°.
5 T.l.c. CH-EA (50:50), Rf 0.77 [alpha]_D = +38.4. IR (nujol) : 3400(NH); 1707 and 1653
10 (C=O), 1597 and 1551 (C=C) cm⁻¹. ¹H-NMR: 7.48(d); 7.46-7.20(m); 7.16(m);
7.04-6.94(m); 6.92(s); 6.37(d); 5.29(d); 4.48(d); 3.38(d); 1.85(m); 1.64-1.30(m).

EXAMPLE 46(+)-N-[1-(Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-ethoxycarbonylphenyl)urea

3-Ethoxycarbonylphenyl isocyante (0.152ml) was added to a solution of intermediate 97 (0.490g) in dry acetonitrile (20ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 1h, then diluted with dichloromethane (20ml) concentrated under vacuum and the residue was triturated with diethyl ether to give the title compound as a white solid (0.543g). M.p. 220-1°. [alpha]_D = +60.8, (CHCl₃, c=1.020) T.l.c. CH-EA (2:1), Rf 0.35. IR: 1709, 1670 and 1690 (C=O)cm⁻¹; ¹H-NMR: 7.93(t); 7.64(m); 7.44-7.39(m); 7.35-7.27(m); 7.24-7.14(m); 7.38(bs); 6.89(dd); 6.58(d); 5.31(d); 4.50(d); 4.34(q); 3.38(d); 1.85(m); 1.61(m); 1.51(m); 1.45(m); 1.37(m); 1.35(t).

25 EXAMPLE 47(+)-N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5benzodiazepin-3-yl]-N'(3-carboxyphenyl)urea

Aluminium iodide (0.137g) was added to a suspension of Example 46 (0.10g) in dry acetonitrile (10ml). The reaction mixture was stirred 6h at 80° then cooled to 23°, diluted with dichloromethane (30ml) and poured into ice (10g). The aqueous layer was acidified
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with a 10% solution of hydrochloric acid (1ml), washed with 5% solution of sodium thiosulphate (20ml) and extracted with dichloromethane (2x25ml). The collected organic phases were washed with water (30ml) and brine (10ml) dried and evaporated to give a white solid (0.118g). This material was purified on silica gel, eluted with CH/EA 1/1 and 5 then EA/Methanol 1/1 to give the title compound (41mg). T.l.c. EA, Rf 0.64. IR: 3354 (NH and OH), 1701 and 1684 (C=O) cm⁻¹; ¹H-NMR: 9.21(s); 7.9-7.8(m); 7.6-7.16(m); 7.0-6.9(m); 4.99(d); 4.30(d); 3.60(d); 1.83(s); 1.65-1.2(m).

EXAMPLE 48

10 N-[2,4-dioxo-7-fluoro-1-(3-methylbut-1-yl)-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]N'-(3-dimethylamino)phenylurea

3-Dimethylaminophenyl isocyante (0.055g) was added to a solution of the intermediate 100 (0.08g) in dry acetonitrile (5ml) under a nitrogen atmosphere. The mixture was stirred at 23° for 30min.; the obtained precipitate was filtered and washed with ethyl ether 15 to give the title compound as a white solid (0.086g). M.p. 249-51°. T.l.c. CH-EA (1:1), Rf 0.5. IR: 1705, 1672 and 1636 (C=O), 1607 (C=C) cm⁻¹; ¹H-NMR: 7.0-7.50(m); 6.81(bt); 6.68(dd); 6.62-6.46(dd); 6.51(bs); 6.28(d); 5.31(d); 4.54-3.60(m); 2.92(s); 1.6-1.40(m); 0.95-0.85(d).

Pharmacy ExampleCapsules or Tablets

	mg/dosage form
5	
Active ingredient	0.1
Polyethyleneglycol	15.0
Lactose	52.4
Starch	30.0
10	
Magnesium stearate	0.5
Silicon dioxide	1.0
Sodium Lauryl Sulphate	1.0
	<hr/>
	100.0

15

The active ingredient is dispersed in a suitable solvent (e.g. ethanol) together with polyethyleneglycol. The solvent is removed. The powder so obtained is blended with the other excipients. The blend can be used to fill gelatine capsules or compressed using appropriate punches. The tablets can be coated using conventional techniques and 20 coatings.

25	Active ingredient	0.1
	Povidone	15.4
	Lactose	74.0
	Hydrogenated vegetable oils	3.0
	Silicon dioxide	1.0
	Sodium Laauryl sulphate	1.5
	Crospovidone	5.0
		<hr/>

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100.0

The active ingredient is dispersed in a suitable solvent (e.g. ethanol) together with povidone. The solution is sprayed on to lactose and the solvent removed. The powder obtained is blended with the other excipients. The blend is used to fill gelatine capsules or 5 compressed using appropriate punches. The tablet can be coated using conventional techniques and coatings.

Oral liquid

Active ingredient	70-100 micrograms/dose
ethanol	5-15%
Sodium saccharinate	0.1-1%
Propylene glycol	q.b. 100%

Injection Formulation

Active ingredient	0.1-100 microgramms
Sodium phosphate	1.50 mg/ml
NaOH	qs desired pH (range 3-9)
glycerol	10-500 mg/ml
water for injection	qs to 0.5-10ml

20

Pack in glass (ampules) with a rubber stopper (vials, syringes) and a plastic/metal overseal (vials only). An inert gas atmosphere (for example nitrogen) may be introduced into dead space of container.

25 CCK - Antagonist Activity

The CCK-A antagonist and CCK-B antagonist activities of compounds of the invention were determined using the guinea pig isolated ileum longitudinal muscle myenteric plexus preparation. The compounds were tested using the procedure G Dal

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Forno et al J. Pharmacol. Exp & Ther. 261 - 1056-1063 1992 and the pK_b value for each compound was determined.

The results obtained with representative compounds of the invention were as follows:

	Compounds of Ex No.	pK _b	
		CCK-A	CCK-B
5	4	5.5	10.1
	14	6.8	8.5
	25	<6.0	9.1
	26	≤5.5	8.7
	45	5.9	8.9

CCK - Receptor Binding

15 The binding affinity of the compounds of the invention for the CCK-A receptor (Pancreas Assay) and CCK-B receptor (guinea pig cortex assay) was determined using the procedure of G Dal Forno et al J. Pharmacol. Exp & Ther. 261 - 1056-1063. The pKi values determined with representative compounds of invention were as follows:

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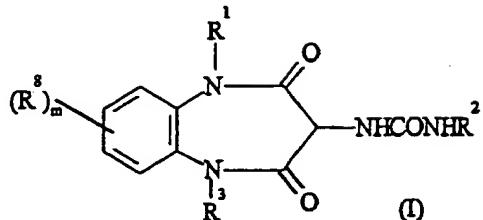
	Compound Ex No	pKi	
		CCK-A	CCK-B
5	4	6.9	9.6
	7A	6.33	8.71
	9	6.02	8.31
	11	5.80	8.01
	14	6.15	8.64
	16	6.95	9.17
10	25	6.49	8.81
	26	6.30	8.81
	27	6.83	9.54
	30	7.00	9.14
	31	6.76	8.82
	39	6.52	8.72
15	41	6.09	8.53
	45	5.95	9.02

The compounds of the invention are essentially non-toxic and therapeutically useful doses. Thus fore example no untoward effects were obserbved when the compound of Example 45 was given orally to mice and rats at doses at which the compound exhibits anxiolytic activity.

CLAIMS

1. Compounds of general formula (I)

5



wherein

- 10 R¹ represents a phenyl, C₃₋₇cycloalkyl, C₇₋₁₁ bridgedcycloalkyl or C₁₋₆alkyl group which alkyl group may be substituted by a hydroxy, phenyl, C₁₋₆alkoxycarbonyl, C₃₋₇cycloalkyl, or C₇₋₁₁ bridgedcycloalkyl group;
- 15 R² represents a substituted or unsubstituted phenyl group (wherein the substitutents may be 1 or 2 of halo, C₁₋₄alkyl, nitro, cyano, trifluoromethyl, trifluoromethoxy, C₁₋₄alkylthio or (CH₂)_n R⁴ wherein R⁴ is hydroxy, C₁₋₄alkoxy, CO₂R⁵ or NR⁶R⁷.
- 20 R³ is phenyl optionally substituted by one or two halogen atoms;
- R⁵ represents hydrogen or a C₁₋₄alkyl group;
- R⁶ and R⁷ independently represent hydrogen or a C₁₋₄alkyl group.
- R⁸ represents hydrogen or a halogen atom; m is zero, 1 or 2;
- 25 n is zero or 1; and pharmaceutically acceptable salts and solvates thereof.

2. Compounds as claimed in Claim 1 wherein R¹ represents phenyl, phenethyl, bridged C₇₋₁₀cycloalkyl, C₄₋₆alkyl, C₃₋₆ hydroxyalkyl, C₁₋₂alkyl substituted by bridged C₇₋₁₀cycloalkyl, alkoxy carbonylmethyl or C₂₋₃alkyl substituted by C₃₋₇cycloalkyl.

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3. Compounds as claimed in Claim 1 or Claim 2 wherein R¹ represents 3-methylbutyl, 3,3-dimethylbutyl, 2-hydroxy-3-methylbutyl, 2-hydroxy-3,3-dimethylbutyl, 2-cyclopentylethyl, 5-norbornenylmethyl or 1-adamantylmethyl.

4. Compounds as claimed in any of Claims 1 to 3 wherein R² represents phenyl optionally substituted by bromine, chlorine, fluorine, methyl, methoxy, methylthio, trifluoromethoxy, cyano, dimethylamino, or (CH₂)_nCO₂R⁵ wherein R⁵ is hydrogen or C₁₋₄alkyl.

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5. Compounds as claimed in any of claims 1 to 4 wherein R² represents phenyl optionally substituted by methoxy, cyano, nitro, methylthio, dimethylamino, ethoxycarbonyl or carboxyl.

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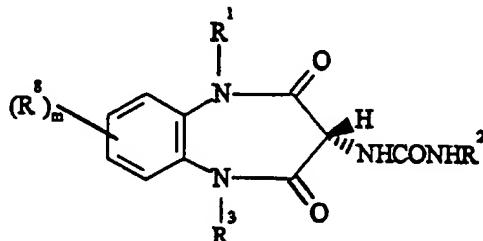
6. Compounds as claimed in any of Claims 1 to 5 wherein R³ represents phenyl optionally substituted by fluorine in ortho or para position.

7. Compounds as claimed in any of Claims 1 to 6 wherein R⁸ represents hydrogen, chlorine or fluorine.

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8. Compounds as claimed in any of Claims 1 to 7 having the configuration

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(1a)

9. N-phenyl-N'-[2,3,4,5-tetrahydro-2,4-dioxo-1-(1-adamantylmethyl)-5-phenyl-1H-1,5-benzodiazepin-3-yl]urea and the (+) enantiomer thereof and salts thereof.

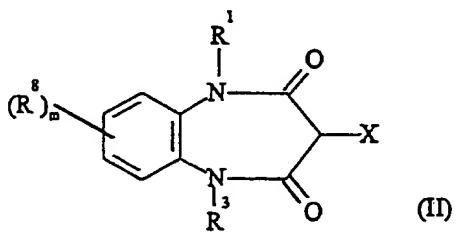
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10. N-[1-(1-Adamantylmethyl)-2,4-dioxo-5-phenyl-2,3,4,5-tetrahydro-1H-1,5-benzodiazepin-3-yl]-N'-(3-carboxyphenyl)urea
N-phenyl-N'-[2,3,4,5-tetrahydro-2,4-dioxo-1-(3-methylbutyl)-5-phenyl-1H-1,5-benzodiazepin-3-yl]urea;

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N-(3-dimethylaminophenyl)-N'-[2,3,4,5-tetrahydro-2,4-dioxo-1-(3-methylbutyl)-5-(2-fluorophenyl)-1H-1,5-benzodiazepin-3-yl]urea; and the (+) enantiomers thereof and salts thereof.

- 5 11. Compounds as claimed in any of claims 1 to 10 for in therapy.
12. The use of a compound as claimed in any one of Claims 1 to 10 in the manufacture of a medicament for the treatment of conditions where a modification of the effects of gastrin and or CCK is of therapeutic benefit.
- 10 13. Pharmaceutical compositions comprising a compound as claimed in any of Claims 1 to 10 in admixture with one or more physiologically acceptable carriers or excipients.
14. A method of treatment of a mammal including man for conditions where
15 modification of the effects of gastrin and or CCK is a therapeutic benefit comprising administration of an effective amount of a compound as claimed in any of claims 1 to 10.
- 15 15. A process for the preparation of compounds as defined in Claim 1 which comprises
20 (a) reacting a compound of formula (II), wherein R¹, R³, R⁸ and m are as defined in formula (I) and X represents the group -N=C=O or NHCOR⁹ wherein R⁹ is an optionally substituted phenoxy group or a 1-imidazole group

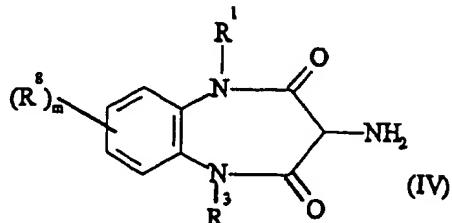


with an amine of formula (III)

wherein R² has the meaning defined in formula (I) or is a group convertible thereto.

(b) acylating an amine of formula (IV)

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10 by reaction with an isocyanate of formula (V) or a carbamoyl chloride of formula (VI).



15 (c) hydrolysis of a compound of formula (I) in which R² is a phenyl group substituted by an alkoxy carbonyl group to yield a compound of formula (I) in which R² is a phenyl group substituted by carboxyl;

and thereafter, if necessary or desired converting the resultant compound, either before or after any separation into its stereochemical isomers into another compound of the invention.

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 93/00098

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all)⁶

According to International Patent Classification (IPC) or to both National Classification and IPC

Int.Cl. 5 C07D243/12

II. FIELDS SEARCHED

Minimum Documentation Searched⁷

Classification System	Classification Symbols
Int.Cl. 5	C07D

Documentation Searched other than Minimum Documentation
to the Extent that such Documents are Included in the Fields Searched⁸III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹

Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
A	EP,A,0 376 849 (ROUSSEL-UCLAF) 4 July 1990 see the whole document -----	1,8, 11-15

⁶ Special categories of cited documents :¹⁰

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

⁷ "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention⁸ "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step⁹ "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art¹⁰ "&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search

16 APRIL 1993

Date of Mailing of this International Search Report

21.04.93

International Searching Authority

EURPEAN PATENT OFFICE

Signature of Authorized Officer

FRANCOIS J.C.

ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.

EP 9300098
SA 69771

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information. 16/04/93

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
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		US-A-	4988692	29-01-91